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UNITED STATES DEPARTMENT OF AGRICULTURE

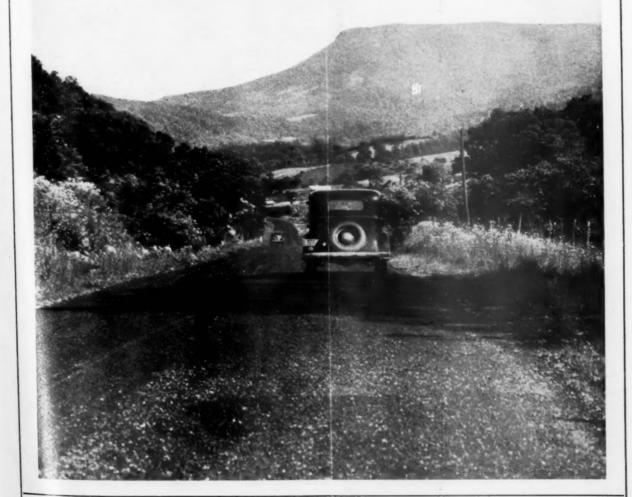
BUREAU OF PUBLIC ROADS



VOL. 16, NO. 6

AUGUST 1935

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### PUBLIC ROADS \*\*\* A Journal of Highway Research

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Volume 16, No. 6

August 1935

The reports of research published in this magazine are necessarily qualified by the conditions of the tests from which the data are obtained. Whenever it is deemed possible to do so, generalizations are drawn from the results of the tests; and, unless this is done, the conclusions formulated must be considered as specifically pertinent only to described conditions.

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### FURTHER STUDIES OF LIQUID ASPHALTIC ROAD MATERIALS

BY THE DIVISION OF TESTS, U. S. BUREAU OF PUBLIC ROADS

Reported by R. H. LEWIS Associate Chemist, and W. O'B. HILLMAN, Assistant Highway Engineer

N A REPORT recently published by the Bureau of Public Roads on A Study of Some Liquid Asphaltic Materials of the Slow-Curing Type, it was shown that the action of sunlight, heat, and air on these materials when exposed in relatively thin films produced residues with physical and chemical characteristics differing greatly from those of the residues developed in the usual laboratory heat tests. It was also shown that when these materials were mixed with a standard sand, molded into cylinders by the Hubbard-Field method, and subjected to the same exposure conditions as the thin films, they developed stability, or bonding strength that could not be attributed entirely to the loss of volatile matter.

### MATERIALS STUDIED TYPICAL OF ALL CLASSES OF LIQUID ASPHAL-TIC MATERIALS FROM PRINCIPAL PRODUCING AREAS

The materials used in the earlier investigation were slow-curing liquid asphalts. They were the products of 10 refineries located in the far and middle West. In further studies conducted in 1933, that are the subject of this report, 32 materials typical of the slow-, medium-, and rapid-curing types of liquid asphaltic materials were used. These samples are identified in table 1 and were the products of 25 refineries located in all sections of the Table 2.—Specification requirements for grades of liquid asphaltic country and probably were made from petroleums of widely different bases and by various refining processes.

TABLE 1 .- Products tested

Sample identifi- cation	Labo- ratory num- ber	Type of material	Pro- ducer	Refin- ery	Location of refinery	Remarks
122 3.44.55.66.77.88.99.110.111.122.133.144.55.166.	1351	SC-2 SC-2 SC-2 SC-3 SC-3 SC-3 SC-2 SC-2 SC-2 SC-1 SC-2 SC-1 SC-2 SC-3 SC-2 SC-2	1 2 2 3 3 4 5 6 7 8 9 9 9 9 2 2 2 2	1 2 3 4 5 6 7 7 8 9 10 11 11 11 12 13	Oklahoma Missouri Illinois do Wyoming Arkansas do Oklahoma West Virginia Rhode Island Louisiana do do Indiana Wyoming do	
17 18 19 20 21 22 23 24 25 26 27 28 29 30 30 31 32	15367 14322 2089 3161 1662 2068 1285 145 145 145 147 147 147 147	SC-2 SC-2 SC-2 SC-2 SC-2 SC-2 SC-2 SC-2	10 10 11 12 13 13 7 14 3 2 2 7 7 15 16 6	15 16 17 18 19 19 20 21 22 12 23 24 25 13	Indiana Illinois	Included in 1932 ex- posure. do. do. do. Steam-reduced Cali fornia residual oil

Furol viscosity was below specification limit.

Furol viscosity was above specification limit.
Penetration of distillation residue was below specification limit.

<sup>1</sup> R. H. Lewis and W. O'B. Hillman, Public Roads, June 1934, vol. 15, no. 4.

As shown in table 1, 23 of the materials were of the slow-curing type, of which 4 samples were tested in 1932 and were included in this study for comparative purposes. Four were of the medium-curing type and five were rapid-curing products. They met the provisional specifications, as given in table 2, of the Bureau of Public Roads and the Asphalt Institute, except as noted in table 1. Samples 21 and 22 were the only slow-curing products for which there was any definite information as to origin or method of manufacture. Both of these materials were steam-reduced California residuals without subsequent blending. All of the rapid-curing products were prepared from 85–100 penetration asphalt and solvent naphtha. The composition of sample 27 was unknown but the other medium-curing products, samples 30, 31, and 32, were, respectively, 110-120 penetration asphalt, 94+ asphaltic road oil, and 100-120 penetration asphalt fluxed with a heavy grade of kerosene. These three medium-curing materials, although subjected to all of the laboratory tests, were exposed only under special conditions that are described later in this report.

road materials investigated

	SC-1	SC-2	SC-3	MC-1	M C-2	RC-2
Flash point, °F Furol viscosity at 77° F., seconds		200+	200+	40-150	150+	80+
Furol viscosity at 122° F., seconds Furol viscosity at 140° F., seconds		200-320	150-300		150-250	200-400
Fotal distillate to 437° F., percent by volume.		2-	2-	10-	2-	10+
Total distillate to 600° F., percent by volume		15-	10-	25+	10-20	20+
by volume		25-	20-	50-	27-	35-
Float at 122° F., seconds Penetration at 77° F	50-	25+	25+	70-300	100-300	60-120
Ductility at 77° F., centimeters. Soluble in CS <sub>2</sub> , percent				60+	60+	60+

The test procedure followed that of the 1932 study except that the fixed-carbon test was omitted as the changes in inherent characteristics that occurred under laboratory and exposure conditions appeared to be more strikingly illustrated by the test for solubility in 86° B. naphtha. Two new tests were added. The Oliensis test 2 for heterogeneity was made on the original materials and on all of the residues, except those from the 50-gram oven-loss test and the 10-week exposure test, and the original materials were examined microscopically. The results of the tests on the original materials are given in table 3 and a detailed analysis of the residues obtained in the routine laboratory tests is given in table 4.

 $^2$  A qualitative test for determining the degree of heterogeneity of asphalts. G. L. Oliensis, Proc. A. S. T. M., vol. 33, pp. 715–728.

Table 3 .- Results of tests on original materials

			Fur	-1	CS3	CCI		1 1							Dist	illati	ion					1	Loss	at 325°	F., 5	hours		
77° F.			visco	sity	luble in	oluble in	86° B. naphtha	liensis tet	Micro-		D	istilla volu	ate b	у	F. (by	weight)	ht)	Test		22º F.	5 sec.	50-gra	am st	ample	20-gr	am sa	1	Asphal residu
Specific gravity at 7	Flash point	Float at 77° F.	At 122° F.	At 140° F.	Organic matter insoluble in	Organic matter insoluble in	Insoluble in 86° B.	Characteristic by Oliensis test	scopic smear test	Initial boiling point	At 374° F.	At 437° F.	At 600° F.	At 680° F.	Distillate to 680° weight)	Loss on cooling (by	Total loss (by weight)	Specific gravity at 77° F.	Index of refraction	Float of residue at 122°	Penetration of resi		Float of residue at 122° F.	Penetration of residue at 77° F., 100 g, 5 sec.	Loss		Penetration of residue at 77° F., 100 g, 5 sec.	Time of reduction
0, 965 1, 056 1, 056 1, 056 1, 043 988 977 966 975 1, 031 966 1, 011 1, 066 977 1, 011 95 1, 086 997 1, 097	3   300   3   300   3   300   3   300   3   3	49 21 38 20 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	2777 175 1977 56 1852 4060 277 2119 201 1996 320 320 320 320 320 320 320 320 320 320	1466 1373 1146 166 1777 11	0 0. 19 3 11 1 13 0 4 6 7 . 38 0 0 1 1 0 5 0 6 4 . 09 7 7 5 5 17 16	. 466 . 477 . 100 . 099 . 111	16. 5 13. 7 9. 4 10. 6 6. 7 18. 1 13. 9 7 . 6 10. 7 11. 3 17. 6 11. 6 10. 7 11. 3 17. 6 18. 1 19. 7 10. 7 11. 3 17. 6 18. 1 19. 7 19. 1 19. 1 19	HO SHOO HHO O SHH HHO O O HHO O SHH HHO O O HHO O HHO O O O HHO O O O HHO O O O O O O O O O O O O O O O O O O O	dododododododo	600 602 602 525 525 594 515 620 500 500 665 525 600 667 536 670 536 580 576 580 580 580 580 580 580 580 580 580 580	3.11.	0 13. 5 18. 11.	1. 3 7. 6 1. 3 1. 3 1. 2 2 3 1. 3 1. 3 5 2. 3 1. 3 5 2. 5 2. 5 2. 5 2. 5 2. 5 2. 5 2. 5 2.	8. (3. (4. (2. (4. (4. (4. (4. (4. (4. (4. (4. (4. (4	0 2.4 1.7 1.7 2.5 5 18.3 1.6 0 20.5 5 12.2 20.5 5 12.2 20.5 5 12.2 20.5 1.0 0 10.2 0 10.3 3 16.3 3 16.3 3 7.0 0 5.1 1.5 0 5.3 1.5 1.5 1.5 1.5 1.5 1.5 1.5 1.5 1.5 1.5	2.8	6. 8 5. 7. 4 23. 4 6. 4 6. 4 6. 4 6. 4 6. 4 6. 4 6. 4	. 926 . 886 . 868 . 868 . 837 . 956 . 837 . 821 . 871 . 871	3 1, 4203 2 1, 4320	66 57 56 56 41 42 56 42 56 42 56 56 56 56 56 56 56 56 56 56 56 56 56	76 77 88 81 61	20. 21. 20. 22. 5 19. 2 21. 17. 5 14.	244 35 30 39 36 31 17 18 12 12 12 12 12 12 12 12 12 12 12 12 12	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	22. 22. 27. 20.	57 38 51 51 57 20 37 38 52 37 38 52 37 38 52 37 38 52 37 7 35 52 37 38 52 37 7 35 52 37 38 52 37 7 35 52 37 38 52 37 38 52 37 7 35 52 37 38 52 37 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 38 52 37 37 37 37 37 37 37 37 37 37 37 37 37	86 56 54 4 55 55	47

1H = Heterogeneous; O=Homogeneous; SH=Slightly heterogeneous.

### DISTILLATION CURVES CLEARLY DISTINGUISH SLOW-, MEDIUM-, AND RAPID-CURING MATERIALS

The distillation curves for the various materials, plotted in figure 1, serve to distinguish the three classes of materials according to their curing properties. It will be noted that the initial boiling points of the slow-curing products, samples 1 to 23, inclusive, were all above 450° F. and in only two instances were they less than 500° F. The medium-curing materials, samples 27, 30, 31, and 32, had initial boiling points as low as 360° F. with none above 450° F.; and the rapid-curing products, samples 24, 25, 26, 28, and 29, all had initial boiling points below 370° F.

With all three classes of materials, the percentage of distillate increased as the temperature increased but not at the same rate. For the slow-curing materials the rate continued fairly uniform up to 680° F., but' with the medium- and rapid-curing products it decreased as the temperature approached 680° F. The decrease in rate was more pronounced for the rapid-curing products. The rate of distillation may be illustrated by expressing the amount of distillate off at any temperature in terms of the total distillate recovered at 680° F. For the medium-curing materials the amount at 600° F. was 65 to 85 percent of the total. For rapid-curing materials the amount was 0 to 50 percent at 374°F., 41 to 85 percent at 437° F., and 88 to 94 percent at 600° F. The residues from the distillation of the slow-curing products were all fluid, while those of the medium- and rapid-curing materials were

semisolid.

That portion of the total loss in the distillation test designated as loss on cooling depends upon the amount of material boiling off immediately above 680° F. As would be expected from the slope of the distillation curves, this loss was greater for the slow-curing products than for the medium- and rapid-curing products. It ranged from 1.2 percent to 5.8 percent with an average of 4.5 percent for the slow-curing products, from 2.9 percent to 4.2 percent with an average of 3.4 percent for the medium-curing products, and from 1.5 percent to 3.1 percent with an average of 2.2 percent for the rapid-curing products.

The total volatile matter in the slow-curing materials as determined by the distillation test, including both the distillate recovered and the loss on cooling, ranged from 2.8 percent to 25 percent with an average of 13.8 percent for the group. The average loss on cooling of 4.5 percent actually represented 36 percent of the total volatile matter in the average slow-curing material used in this study. While this loss on cooling may be unimportant in estimating the relative volatility of various liquid asphalts, it must be considered if the results of the distillation test are to be compared directly with the results of other laboratory and exposure tests.

The different classes of materials are also readily identified by the results of the volatilization and asphaltic-residue tests. The slow-curing products lost in the 50-gram volatilization test from 53 to 76 percent, the medium-curing products from 73 to 80 percent, and the rapid-curing products from 92 to 97

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Table 4.—Results of tests on laboratory residues

				Disti	llatio	n								Los	s at 3	25° F	, 5 h	ours							2	Asph	altie i	residu	e		
				Те	ests or	resid	lue					50-g	ram	samı	ole				20-gr	am sa	mple					P	rests o	n resi	idue		
					Due	atil.		14				Tes	sts or	resi	due				Test	s on i	esdue										
tillation test		Pertrat	ion		cer mei per i	at 5 nti- ters	insoluble in CS <sub>2</sub>	insoluble in CCL	B. naphtha			Per			Ducity a cen met per r	ti- ers nin-			77° F., 100 g., c.	insoluble in	insoluble in	3. naphtha		Per trat			Duc ity s cen met per s	at 5 ti- ers nin-	nsoluble in CS <sub>1</sub>	Organic matter insoluble in CCL	
Total loss in distillation test	Float at 122° F.	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77º F.	At 34°-35° F.	Organic matter insoluble	Organic matter insoluble in	Insoluble in 86°	Loss	-	g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°-35° F.	Loss	Float at 122° F.	Penetration at 75 sec.	Organic matter CS2	Organie matter CCL	Insoluble in 86° B. naphtha	Loss	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°-35° F.	Organic matter insoluble in	Organic matter i	
23. 30. 21. 23. 25. 20.	60 43 45 58 21 22 65 62 1 22 1 22 1 2 2 2 2 2 2 2 2 2 2 2 2 2	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	188 7 288 7 185 5 200 2 25 5 41 5 41 5 41	1172 3 116 5 109 9 122 3 121 117 101	110+ 2110+ 110+ 110+ 2110+ 110+ 110+	0.0.4.4.4.6.4.4.4.4.4.4.4.4.4.4.4.4.4.4.	.31	.41 .34 .17 .08 .13 .10 .2 .37 .11 .10 .15 .10 .15 .10 .15 .10 .15 .10 .10 .15 .10 .10 .10 .10 .10 .10 .10 .10 .10 .10	7. 8 24. 1 18. 3 10. 3 12. 7 11. 7 18. 0 14. 0 10. 7 21. 3 7. 2 9. 5 6. 1 9. 6 23. 0 25. 4 21. 0 26. 8 21. 8 29. 0 29. 0	4. 9 6. 0 5. 7 4. 3 5. 5 20. 6 21. 8 20. 7 22. 1 19. 8 21. 1 17. 6 14. 5	35 30 39 36 17 18 25 42 29 42 10 10 31 24 12 26 43 43 43 42 26 43 42 26 43 44 26 43 44 44 44 46 46 46 46 46 46 46 46 46 46	1522 999 1133 85	355 33 30 24	111 126 120 123 124	110+ 110+ 110+ 110+ 110+	9. (4. )	4. 6 4. 5 8. 6 9. 14. 3 3. 3 6. 9 15. 2 10. 2 8. 6 3. 8 14. 1 15. 6 7. 7 8. 3 10. 7 7. 7 10. 2 2 2 2 3 2 2 2 3 3 1 3 1 3 1 3 3 3 4 3 1 3 1 3 1 3 1 3	277 578 388 529 488 588 588 588 588 588 588 588 588 588	80 56 55 53 411 50 50 50 50 50 50 50 50 50 50 50 50 50	1.17	3.51 1.91 1.14 1.15 1.06 6.5.31 1.44 1.10 1.15 1.55 1.14 1.10 1.15 1.25 1.22 1.23 1.23 1.24 1.23 1.24 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.24 1.27 1.28 1.28 1.28 1.28 1.28 1.28 1.28 1.28	15. 2 10. 0 21. 7 19. 8 12. 2 13. 4 14. 1 20. 9 15. 2 12. 1 23. 3 8. 8 31. 4 11. 1 11. 2 25. 5 28. 4 24. 6 27. 5 24. 9 32. 8 29. 5	28. 3 25. 3 4 7 3 7. 6 35. 2 2 34. 7 3 34. 1 46. 8 38. 2 30. 5 38. 3 3 2 24. 1 37. 4 29. 4 1 37. 4 29. 4 1 22. 8 21. 8 21. 6 21. 5 24. 2 2 24. 2 2 2 2 2 2 2 2 2 2 2 2 2 2	105 107 107 108 99 92 100 86 106 84 93 103 103 103 113 98 101 113 99 101 114 95 101 101 101 101 101 101 101 101 101 10	377 155 168 288 344 117 355 27 36 15 27 111 211 22 35 36 15 27 111 211 22 35 36 36 41 17 27 112 27 27 113 28 36 36 37 41 115 27 27 27 27 27 27 27 27 27 27 27 27 27	109 107 108 113 116 118 114 110 117 113 119 115 117 114 119 119 110 115 120 118 120 118 120 118	Cm 40 110+110+110+110+110+110+110+110+110+11	.0 4.55 5.0 6.0 .1 6.51.0 .0 4.55 5.3 .0 4.8 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0	.555	1. 50 1. 62 12 14 19 15 3. 81 12 03 03 03 3. 41 15 3. 84 04 04 05 05 05 06 07 07 07 08 09 09 09 09 09 09 00 00 00 00	5 0 2 2 2 2 4 9 5 1 2 3 3 3 1 5 5 5 6 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9

tion test. The residues from both volatilization tests of the slow-curing products were fluid; those of the medium-curing products were fluid in the 50-gram volatilization test and semisolid in the 20-gram volatilization test; and those of the rapid-curing products were both semisolid. For the slow-curing products, the loss in neither volatilization test amounted to as much as that in the asphaltic residue test. For the medium-curing products, the loss in the 20-gram volatilization test approximated the loss in the asphaltic-residue test, while for the rapid-curing products the losses in all tests, including the distillation test, were approximately the same. The loss in the 20-gram volatilization test, while always greater than that in the 50-gram volatilization test, was always less than in the distillation test. The residue from the 20-gram volatilization test may, however, be harder or softer than the residue from distillation.

The slow-curing products were reduced to a residue of 100 penetration in from 30 to 420 minutes, with an average of 126 minutes. In producing residues of the same consistency medium-curing products took from 16 to 30 minutes, with an average of 23 minutes and the rapid-curing products took from 11 to 15 minutes, with an average of 13 minutes. When making the asphaltic-residue test on the rapid-curing products, diffi-

percent as much as they lost in the 20-gram volatiliza- not too hard, since the high volatility of the solvent caused the cut-back materials to be reduced to 100 penetration before the temperature for making the test was reached.

### MAJORITY OF MATERIALS SHOWED A CLEAR FIELD ON MICRO-SCOPIC EXAMINATION

The microscopic test, adopted by one State highway department for the detection of cracking-coil products, was made on all materials. The specification of the State did not set up an exact procedure. It merely stated that a freshly prepared smear of the asphaltic material diluted with carbon tetrachloride should show a clear field free from carbonaceous matter when subjected to a magnification of 200 diameters. In the work covered by this report, the test was standardized by using 2 parts by weight of carbon tetrachloride and 1 part by weight of asphaltic material in preparing the slides for observation. When prepared in this manner all of the materials but seven showed a clear field. In preparing the slides for the photomicrographs illustrated in figure 2, 6 parts by weight of carbon tetrachloride and 1 of the asphaltic material were used.

Because carbon tetrachloride has both solvent and flocculent properties, its use as a diluent was questioned. Therefore, slides were also prepared with the undiluted materials and it was found that only those culty was experienced in obtaining a residue that was materials that showed flecks when undiluted showed

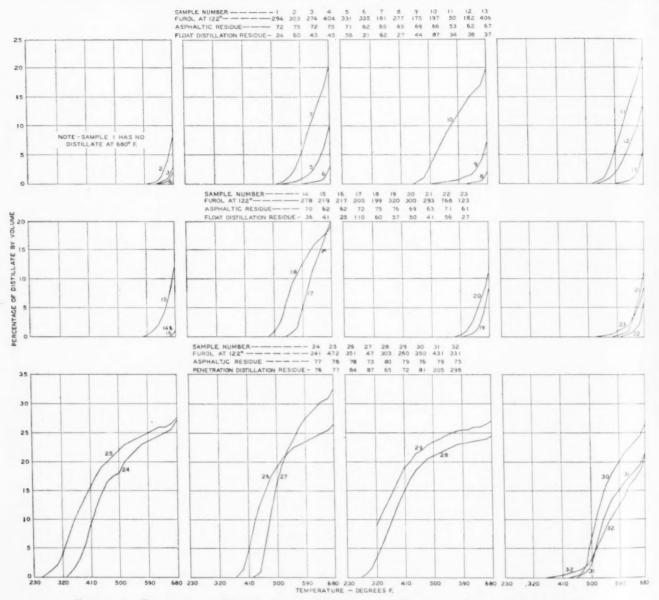


FIGURE 1.—RELATION OF PERCENTAGE OF DISTILLATE BY VOLUME TO DISTILLING TEMPERATURE.

them in the 2 to 1 and 6 to 1 dilutions. This indi- probable that the flecks shown are particles of free cates that the insoluble matter was already flocculated and that carbon tetrachloride, in the quantities used, did not precipitate carbonaceous flecks in those materials not containing them when undiluted. Recently a sample was tested that contained flecks when undiluted that disappeared on dilution with carbon tetrachloride; and another sample that was practically clear when undiluted contained flecks when diluted. In the first case, the poor solvent properties of the distillate used in the material were responsible for incomplete solution of the base asphalt, that immediately went into solution with the addition of carbon tetrachloride. In the second case it was quite evident that the carbon tetrachloride acted as a flocculent. However, since 6 of the 7 samples that contained carbonaceous material had relatively high percentages of material insoluble in carbon tetrachloride, it is extending from the top of the box downward at an

carbon and carbenes.

In conducting the exposure tests three samples of each material were placed in seamless, flat-bottom tins having a diameter of 51/2 inches and a depth of five-eighths inch. Fifty cubic centimeters of material were used to obtain a uniform film or layer thickness of about one-eighth inch. The samples were then placed in exposure boxes made of wood. A plateglass cover resting on strips of felt fastened to the edges of each box made a tight joint and excluded all dust and dirt. A current of air was passed through a wash bottle containing sulphuric acid to remove dust and eliminate moisture, and was admitted through the bottom of the boxes and escaped through slots in the sides, thus serving to carry off the vapors formed. The slots were protected from rain by this boards

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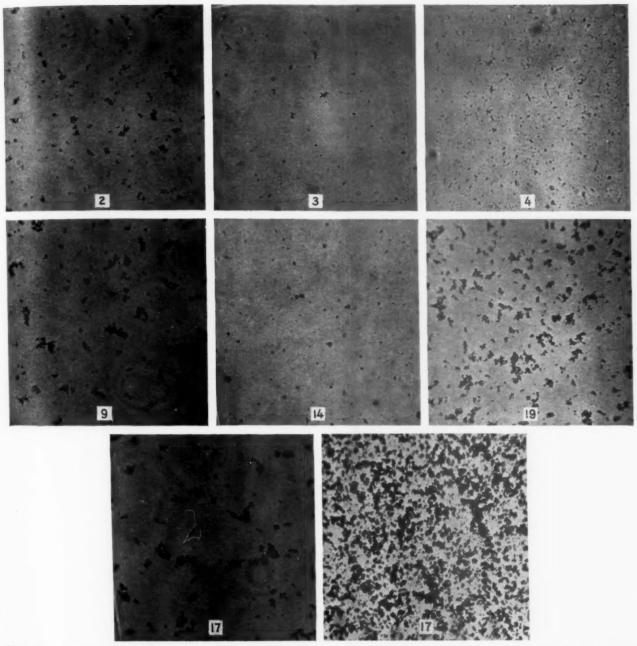


Figure 2.—Photomicrographs of Materials Containing Carbonaceous Matter (Magnified 200 Times). The Lower Two Illustrations Show Results Obtained With Sample 17; in the One on the Left the Material Was Diluted with Carbon Tetrachloride and the One on the Right Shows Undiluted Material.

provided a means of determining the temperature. The assembly of the boxes is shown in figure 3.

### DIFFERENCES FOUND BETWEEN SLOW-CURING AND CUT-BACK PRODUCTS AFTER EXPOSURE

The samples were placed in the boxes on June 15, 1933, and were weighed periodically to determine the loss in weight. A complete set of samples was removed

angle of about 45°. Cotton batting inserted in the heat of the sun and varied with the amount of sunshine. slots excluded dust. A thermometer in each box On clear days the temperature was extremely high, the On clear days the temperature was extremely high, the maximum recorded being 196° F., but on days with no sunshine the temperature in the boxes was the same as that of the air. During the period of exposure the average maximum daily air temperature was 85° F. The samples exposed for 5 weeks were subjected to 333 hours of sunlight and those exposed for 10 and 15 weeks were subjected to 611 and 866 hours of sunlight, respectively. The percentage of loss at different periods of exposure and tested at the end of 5 weeks, another set at 10 is given in table 5 and the results of tests on the weeks, and the last set at 15 weeks. The temperature residues are given in table 6. Photographs of typical of the boxes was dependent entirely upon the radiant surfaces at the end of 15 weeks are shown in figure 4.

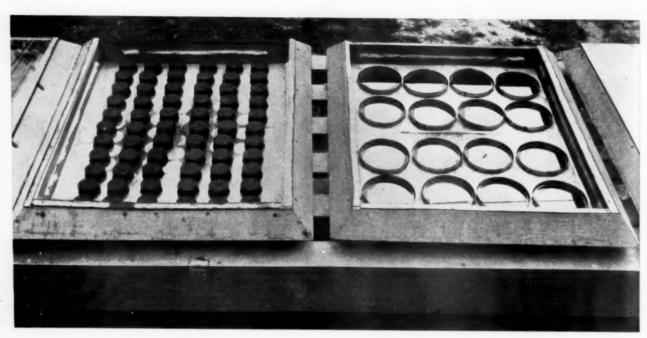


FIGURE 3.—STABILITY SPECIMENS AND THIN FILMS OF MATERIAL EXPOSED TO SUNLIGHT.

While the majority of the materials progressively lost weight during the period of exposure, a number of them actually gained at first, although they later lost more than the amounts gained. An exception to this was sample 1, which had gained 3.6 percent at the end of 8 days and at the end of 15 weeks still showed a slight gain. The samples exposed for 15 weeks were used in determining the loss at 2, 8, 15, 22, 50, and 105 days, while the percentage of loss given for the 35- and 70-day exposures was based on samples used for test at the end of 5 and 10 weeks, respectively. This was done to eliminate errors in calculating the results of subsequent tests made upon the respective residues and accounts for slight variations that may appear to indicate gains instead of losses.

As was expected, the cut-back products lost weight very rapidly. At the end of 2 days they had lost from 86 to 92 percent of their maximum loss with an average of 89 percent, and at the end of 35 days they had lost from 97 to 100 percent with an average of 99 percent. For the slow-curing products the rate of loss was much less, but was considerably more variable. In 2 days those samples that had undergone a loss had lost from 3 to 60 percent of their maximum loss with an average of 35 percent. In 15 days they had lost from 16 to 84 percent with an average of 50 percent, in 35 days from 63 to 100 percent with an average of 82 percent, and in 70 days from 74 to 100 percent with an average of 89 percent. Some idea of the relative speed of curing or volatility of the two types of material may be obtained by comparing their losses under these exposure conditions. The slow-curing materials took 70 days to lose an average of 89 percent of their volatile matter, while the cut-back materials underwent the same percentage loss in 2 days.

### LOSS IN DISTILLATION TEST MOST NEARLY APPROXIMATED LOSS IN 15 WEEKS' EXPOSURE FOR ALL TYPES OF MATERIALS

Figure 5 shows the relation between the percentage of loss upon exposure and loss in the distillation test, terials studied, both in 1932 and 1933, the total loss in

TABLE 5 .- Loss in thin film exposure

			Lo	oss on exp	posure fo	r—		
Sample identi- fication	2 days	8 days	15 days	22 days	35 days	50 days	70 days	105 days
	Percent	Percent	Percent	Percent	Percent	Percent	Percent	Percen
	-2.6	-3.6	-3.4	-3.4	-3.0	-2.7	-2.7	-0.
	1.9	4.3	5. 7	7.6	9.5	10.3	10.2	12.
	2	2.0	2.9	4.7	6.6	8.0	8.0	9.
		1.5	1.7	.3.2	5. 1	4. 4	5.8	5.
	2.6	5.4	6.5	8.2	9.5	9.8	9.9	11.
	3	1.2	2.0	4.1	5. 1	6. 1	8.5	8.
	7.8	12.9	14.6	17.7	20.2	20.0	21.0	21
	-1.8				2.3		1.8	2
	. 3	2.4	4.0	5.3	8. 1	8.2	9.6	10
0	11.0	14.4	14.5	15. 7	16.4	16.7	16. 8	17
1	7.8	14.7	16, 4	18. 2	20.4	21.0	21.6	22
2	4.6	8.9	9.4	11.8	13.7	13.4	13.8	15
3	.4	2.6	4.3	6. I	7.3	10.1	8.5	9
4	-1.3	. 2	1.3	3.4	5. 2	6.9	6.0	8
5	1.9	5.5	6.9	8.4	9.3	12. 1	10.6	13
6	-3.5	-1.5	-1.0	1.8	4.6	5. 7	5.7	6
7	8.6	12.6	14.4	15. 9	18. 5	18.9	18.7	19
8	11.5	15.7	16.5	17.4	18. 2	19. 2	17.7	19
19		5.0	6.1	8.6	10.1	12.4	12, 0	13
20		6.8	7.8	9. 8	11.7	12.6	11.5	13
21	-1.1	3.7	6.4	8. 2	12.5	13.6	12.9	14
22		3.4	5. 1	6. 7	8.5	9.8	9.1	10
3	1.3	3.7	4, 6	6. 6	8.3	10.6	13. 1	11
24		21. 2	21.5	22.0	22. 2	22. 2	22.5	22 22
25		21.6	21.9	22. 1	22. 2	22.0	22. 0	22
26		21.6	21.8	22. 1	22. 4	22. 1	22, 1	27
27		27.1	27.3	27.5	27. 6	27.6	28.3	20
28 29		19.6 20.8	19.9	20. 0	19. 9 21. 6	20. 1	20.0	21

loss in the two oven tests, and loss in the asphaltic-residue test. Figure 5 indicates that for the slow-curing products the loss in 15 weeks' exposure was about 2½ times as great as that in the oven test on a 50-gram sample, about 1½ times as great as that in the oven test on a 20-gram sample, and about the same as the loss in the distillation test. No relationship was apparent between the loss in the exposure and asphaltic-residue tests. The loss in the latter test was, however, invariably greater, ranging from 1½ to 14 times the loss occurring in 15 weeks' exposure with an average of 2½ times this loss. For the cut-back products the loss in all tests was approximately the same. In all the materials studied, both in 1932 and 1933, the total loss in

FIGURANTES

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Table 6.—Results of tests on exposure residues

			5-W	ek e	xposu	ге							10-w	eek e	xposu	re							15-we	ek ez	xposui	ге		
		Pene			Ducti at 5 c time per n ut	ters	ble in C83	ble in CCL	naphtha			Pen trati			Ducti at 5 c time per n ut	ters	ible in CS2	ible in CCl,	naphtha			Pen			Ducti at 5 c timet per m ute	en- iers	uble in CS2	ible in CCI,
	Float at 122° F.	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°-35° F.	Organic matter insoluble	Organic matter insoluble in	Insoluble in 86° B. nap	Loss	Float at 122 ° F.	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°-35° F.	Organic matter insoluble	Organic matter insoluble	Insoluble in 86° B. nap	Loss	Float at 122° F.	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°-35° F.	Organic matter insoluble	Organic matter insoluble
-2. 6 9. 5 6. 6 5. 1 9. 5 -0. 3 20. 2 2. 3 8. 1 1, 1 10. 4 13. 7 7, 3 5, 2 9, 3 4. 6 18. 2 10. 1 11. 7 12. 5 8, 5 8, 1 11. 7 18. 2 19. 5 19. 5 1	Sec. 68 153 200 34 123 163 000+ 131 110 129 105 150 70 197	26	39 80 22 28 9 14 14 7 7	966 110 106 1366 93 98 98 98 1146 155 151 138	95. 0 95. 0 17. 0 5. 0 4. 5 8 4. 0 7. 0	5.00 1.00	2. 93 1. 28 3. 22 1. 63 1. 81 4. 60	25 7. 51 5. 69 7. 57 11. 69 1. 15 20 2. 21 2. 21 2. 21 2. 21	25. 6 25. 9 15. 3 21. 0 18. 7 26. 4 26. 6 23. 8 21. 0 21. 8 29. 4 25. 9 19. 9 30. 8 14. 8 34. 4 21. 4 18. 2 19. 1 24. 7 32. 4 31. 7 27. 9 27. 9	1. 8 9. 6 16. 8 21. 6 13. 8 8. 5 6. 0 10. 6 5. 7 17. 7 12. 0 11. 5 12. 5 9. 1 13. 1 22. 5 22. 0 22. 1 28. 3 20. 0	44 163 356 1,000 +	97 300+	53 35 31 47 14 43 10 41 14 10 13 13 6 9	127 134 118 93 129 129 121 105 119 103 117 101 105 121 112 95 104 147 161 158 163 158	3.5 1.0 10.0 17.0 40.0 105.0 53.0 .5 110+ 110+ 90.0 110+ 1.0 5.5 4.8 3.0 4.5	.00 3.88 2.33 .55 3.55 1.5 .00 7.55 .00 3.88 7.33 .00 .00	3. 08 3. 17 6. 50 4. 43 3. 34 4. 43	13. 34 11. 29 7. 41 	31. 3 29. 3 30. 5 17. 0 22. 1 22. 5 30. 2 29. 4 29. 3 23. 9 27. 3 31. 5 30. 3 23. 7 24. 2 22. 3 28. 2 32. 8 36. 9 31. 5	9. 4 5. 8 11. 8 2 21. 8 2 21. 8 2 21. 8 2 10. 7 17. 3 15. 2 9. 8 8. 1 19. 6 14. 3 10. 1 11. 1 11. 1 12. 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	7 1,000+7 3 3 3 4 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	197 20 49 49 59 52 195 76 51 77 280 64 245 29 93 29 41 166 112 49 13 24 17 17	18 23 76 104 60 477 222 844 83 321 30 999 122 82 22 6 6 13 13 5 8	154 144 153 135 107 133 156 135 126 112 133 109 168 120 103 111 129 160 163 161 153	20. 0 .3 .8 .5 .5 .21. 0 .55. 0 .55. 0 .7. 5 .9. 0 .41. 0 .47. 0 .47. 0 .50. 110+ .110+ .70. 0 .50. 0 .50	.0 .0 .0 .5 .5 .0 .0 .3 .0 .0 .4 .3 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0 .0	5.71 5.69 3.36 45 5.48 3.07 7.50 	112. 088. 400. 177. 178. 188. 418. 188. 188. 188. 188. 188. 18

1 The 10- and 15-week exposure residues of this sample were nonhomogeneous.

<sup>3</sup> None of the residues after exposure of this sample was homogeneous.

the distillation test most nearly approximated the loss in 15 weeks' exposure.

At the end of 15 weeks' exposure the surfaces of the samples varied greatly in appearance, as shown by the typical photographs in figure 4. The appearance of sample 14 was typical of samples 1, 2, 3, 4, 5, 6, 7, 9, 12, 18, 19, 20, 21, and 22. However, there were some variations in actual appearance that could not be shown in a photograph. Samples 1, 2, and 3 had mottled, slightly greasy surfaces. Samples 4, 9, and 19 had mottled, slightly greasy surfaces that were slightly iridescent. Samples 14, 21, and 22 had mottled, iridescent surfaces that were not greasy. Sample 5 had a uniformly mottled appearance neither iridescent nor greasy. Samples 6, 7, 12, 18, and 20 were smooth and glossy.

Sample 23 was mottled and slightly greasy, like samples 1, 2, and 3, but had some dull areas as shown. Sample 13, while smooth and glossy, had some dull areas as shown; sample 16 was similar although it was slightly mottled.

Samples 10, 11, and 15 were checked over most of the area of their surfaces; their condition is shown by the photograph of sample 15. The unchecked areas in samples 10 and 11 were glossy but in sample 15 the surface was neither dull nor glossy. Samples 8 and 17 had very rough surfaces, shrunken and pitted as shown by their photographs. The material in the bottom of the cracks was soft but the outer surface was very hard. The cut-back products were all very rough and wrinkled, as shown by the photographs of samples 26, 27, and 28. slow-curing materials all developed residues after 15

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The material in the cracks was glossy while the outer surface was dull.

At the end of 15 weeks all of the cut-back products were exceedingly hard. As they were originally combinations of semisolid asphalt and volatile fluxes and became semisolid in the laboratory tests, it would be expected that a semisolid residue would rapidly develop upon exposure. It has been believed generally that the asphalt base used in cut-back materials is relatively resistant to changes caused by weathering and that after the cutting medium is removed the character of the material changes but little. The six cut-back materials used in this study, while losing but little weight after 2 days, were much harder and had much less ductility at the end of 5 weeks' exposure than the asphalt used as base material.

The slow-curing products varied as greatly in consistency as in the rate and amount of volatile matter given off during exposure. At the end of 5 weeks all of them were harder than the residues from the distillation and oven tests, but only 5 had a penetration at 77° F. of less than 200. The consistency varied from a float of 34 seconds at 122° F. to a penetration of 51 at 77° F. At the end of 15 weeks only 3 materials had a penetration at 77° F. of over 200 and one of these materials had disintegrated to such an extent that no true penetration test was possible. The consistency ranged from a float of 71 seconds at 122° F. to a penetration of 20 at 77° F. In the previous work, the

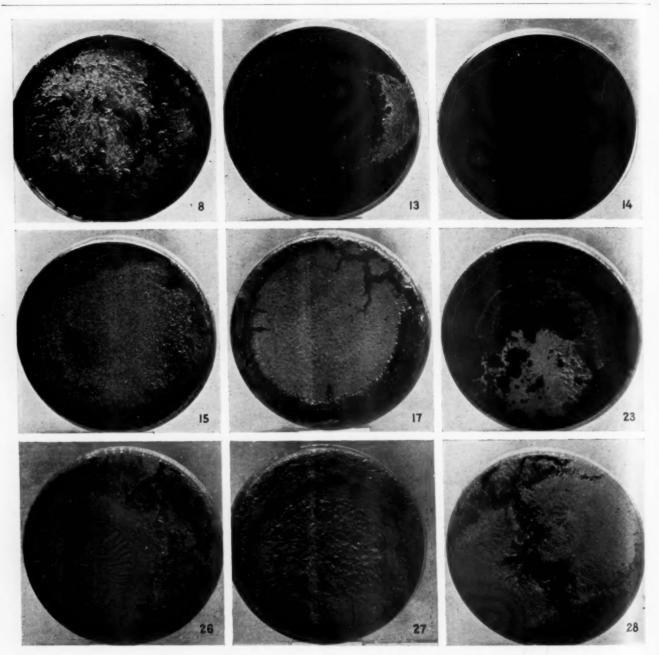


FIGURE 4.—CONDITION OF SURFACES AFTER 15 WEEKS OF EXPOSURE.

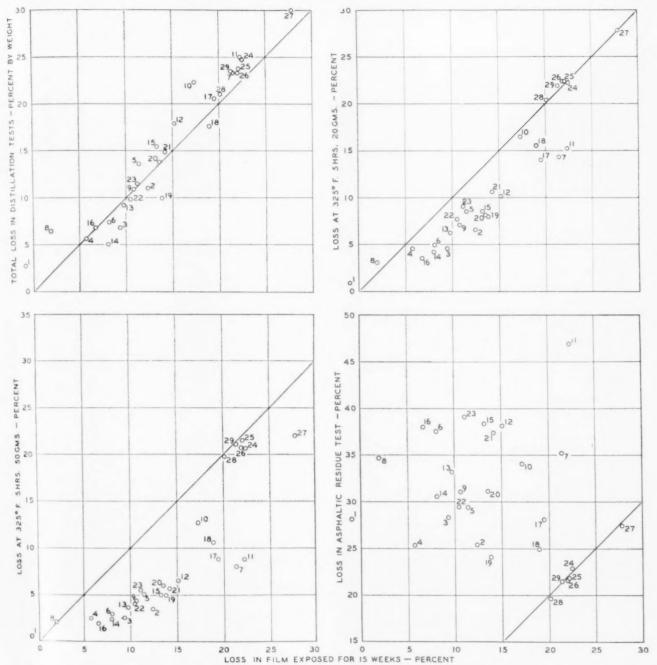
weeks that were as hard as or harder than their asphaltic | timeters and 15 had ductilities at 34°-35° F. of 3 deenresidues of 100 penetration. It seems that it is impossible to predict, from the results of any of the laboratory tests, the consistency of the residues after exposure. Generally, however, those materials that took a long time to be reduced to 100 penetration in the asphalticresidue test and those whose residues from the distillation and oven tests had a low float-test value were the softest or most fluid at the end of 15 weeks.

### ASPHALTIC RESIDUES DIFFERED IN CERTAIN RESPECTS FROM RESIDUES AFTER EXPOSURE OF ABOUT THE SAME PENETRATION

A comparison of the residues after exposure and asphaltic residues is of interest. All but 5 of the asphaltic although having the sa residues had ductilities at 77° F. of more than 110 cenably in other respects.

timeters or more. After 15 weeks' exposure only 7 products had ductilities at 77° F. over 50 and only 4 over 110. Only 6 had ductilities at 34°-35° F. over 3½. These differences in ductilities may, in a number of instances, have been caused by differences in the consistencies of the residues after exposure and asphaltic residues.

However, in some cases one of the residues from exposure had approximately the same penetration as the asphaltic residue and the laboratory residues and residues after exposure may be compared directly. 7 shows the results of tests on both such residues and indicates that the two residues from the same material, although having the same consistency, varied consider-



COMPARISON OF THE PERCENTAGE OF LOSS AFTER 15 WEEKS OF EXPOSURE WITH LOSS IN THE LABORATORY EVAPORATION TESTS.

The ratio of the penetration at 77° F. to that at 32° and carbon disulphide, the percentages were much F. was always lower for the residue from exposure. The ductility at 77° F. of the residue from exposure was less than that of the asphaltic residue in all but two cases, and for these the ductility of both residues In five cases the ductility at 34°-35° F. of the residue from exposure was greater than that of the asphaltic residue. In every case except two the percentage of material insoluble in naphtha was greater for the residue from exposure than for the asphaltic residue. In every case where there was an appreciable

greater for the residues from exposure.

Figure 6 shows the development of free carbon, carbenes, and asphaltenes in laboratory and exposure tests for the samples that originally contained or finally developed carbenes in appreciable amounts. Samples 8, 23, and 27 developed only relatively small amounts of carbenes and the development of the insoluble constituents is shown only for sample 27. In figure 6 the volatile matter and the material insoluble in carbon residue. In every case where there was an appreciable amount of material insoluble in carbon tetrachloride are plotted for the original materials and their residues.

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Table 7.—Comparison of residues of approximately the same penetration from the asphaltic-residue test and from exposure

				Asph	altic resid	lue							Re	sidue fron	n exposu	re			
ation	Penetr	ation	77° F.		Ductilit per m	y,5 cm inute	86° B.	cl, in-	Si In-	Peneti	ration	77º F.		Ductilit per m	y, 5 cm inute	86° B.	ccl,	S <sub>2</sub>	9
Sample identification	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Ratio: Pen. at	Softening point	At 77° F.	At 34°-35° F.	Insoluble in 8 naphtha	Organic matter soluble in CCL	Organic matter soluble in CS	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Ratio: Pen. at	Softening point	At 77° F.	At 34°-35° F.	Insoluble in 8 naphtha	Organic matte soluble in C	Organic matter soluble in CS <sub>2</sub>	Time of exposure
2 3 4 5 10 11 13 14 15 18 20 22	100 105 107 105 106 84 93 93 103 88 102 113	15 15 18 28 35 21 27 13 25 27 21 23	6. 7 7. 0 5. 9 3. 8 3. 0 4. 0 3. 4 7. 2 4. 1 3. 3 4. 9 4. 9	°F. 109 107 108 113 117 113 115 110 115 120 114	Centi- meters 110+ 110+ 110+ 110+ 110+ 110+ 110+ 110	4.5 4.8 .8	Per- cent 35. 2 32. 5 25. 5 24. 2 25. 6 21. 1 22. 7 31. 1 23. 1 18. 5 19. 5 14. 5	Per- cent 2 16 1.50 1.62 .12 .03 .09 .72 .03 .15 .04	Per- cent 0.16 .16 .19	93 84	34 222 33 37, 53 35, 39, 21, 47, 28, 28, 28,	2.5 2.8 3.4 2.6 1.8 2.1 2.3 3.8 2.2 3.3 3.0 4.0	°F. 108 127 134 118 129 121 119 112 117 116 120 111	Centimeters 63.0 5.0 4.3 87.0 10.0 17.0 40.0 97.0 53.0 110+ 72.0	3.0	Per- cent 30, 2 31, 3 29, 3 30, 5 29, 4 29, 3 27, 3 33, 4 30, 3 8 27, 2 24, 2	Per- cent 9. 23 11. 29 7. 41 8. 61		Wee

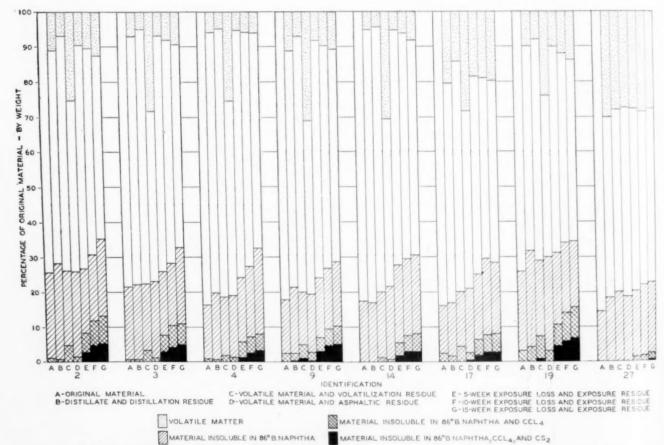


FIGURE 6.—Composition of Selected Materials and Their Residues as Determined by Solubility Tests.

All percentages are expressed in terms of the weight of the original material.

The solid portion in each vertical column represents free carbon or material insoluble in carbon disulphide. The remainder represents bitumen or material soluble in carbon disulphide. The double cross-hatched portion represents carbenes or bitumen insoluble in carbon tetrachloride, and the single and double cross-hatched portions together represent asphaltenes or bitumen

insoluble in 86° B. naphtha. The remainder of the column represents material soluble in 86° B. naphtha that, in the original material, includes the more volatile hydrocarbons vaporized and lost under test conditions, as shown by the dotted area. The materials soluble in 86° B. naphtha have been termed "malthenes" by Richardson.<sup>3</sup> This designation, however, has not been generally accepted in the United States.

<sup>8</sup> Clifford Richardson, The Modern Asphalt Pavement, p. 544 (2d ed.).

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As shown by figure 6, the material insoluble in naphtha included carbenes and free carbon; in fact, in the residue from 15 weeks' exposure of sample 19 the material insoluble in naphtha contained 19 percent free carbon and 27 percent carbenes. The term asphaltenes also includes carbenes. Nellensteyn 4 has stated that asphaltenes, carbenes, and free carbon all consist of the same matter, that is, dispersed carbon in decreasing states of protection. The so-called "protective bodies" he designates as "micelles." He states that when extracted asphaltenes are heated at a temperature of 527° F. they will be changed largely to free carbon, but a normal asphaltic material can be heated at 662° F. for a long period with little production of free carbon. The reason for this is that in normal asphalts the amount of protective bodies, or micelles, is such that the decomposition of part of them influences their protective qualities only slightly. Marcusson 5 states that the material soluble in 86° B. naphtha (malthenes) is composed essentially of oily constituents and asphaltic resins. As the time of exposure increased, the percentage of asphaltenes, carbenes, and free carbon increased while the percentage of malthenes decreased.

In the 1932 investigation only those materials that had high specific gravities and initially contained some material insoluble in carbon disulphide with appreciable amounts insoluble in carbon tetrachloride developed carbenes either in laboratory or exposure tests. In the present study some of the materials such as samples 14 and 27, that originally had high solubilities in carbon disulphide and carbon tetrachloride, developed carbenes even during some of the laboratory tests. In those materials in which carbenes and free carbon were developed, it may be considered that the amount or the protective quality of the micelles was insufficient to

prevent carbonization.

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All of the rapid-curing products at the end of 15 weeks had developed residues containing about 0.5 percent of material insoluble in carbon disulphide and carbon tetrachloride. For all materials the solubility

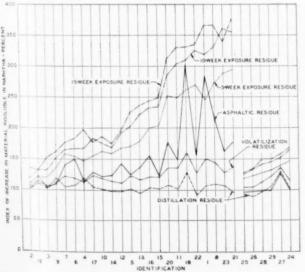


FIGURE 7.—INDEX OF INCREASE IN MATERIAL INSOLUBLE IN NAPHTHA IN THE VARIOUS RESIDUES.

in carbon disulphide was almost the same as the solubility in carbon tetrachloride, indicating the almost complete absence of carbenes.

### MATERIAL INSOLUBLE IN NAPHTHA DEVELOPED MOST UNDER EXPOSURE AND LEAST IN THE DISTILLATION TEST

In figure 6, where all the percentages are expressed in terms of the weight of the original material, if the percentage of material insoluble in naphtha in any residue is divided by the percentage of material insoluble in naphtha in the original material and multiplied by 100, the result is the index of increase in material insoluble in naphtha. An index of 100 therefore indicates no change in the amount of material insoluble in naphtha. This index for all the samples is plotted in figure 7 and shows that generally there was an increase in material insoluble in naphtha during the various tests. Generally, this index was least for the distillation test and greatest for the 15-week exposure test, In the distillation test it ranged from 88 to 129.

While inaccuracies in testing may account for indexes of less than 100 in the distillation residues, it is possible for indexes actually to be below 100. Several samples of very viscous, slow-curing asphaltic material, really semisolid asphalts, were recently subjected to the distillation test. The materials did not yield any distillate but there was considerable loss on cooling. Nevertheless, the residues were softer than the original materials and contained less material insoluble in

naphtha.

In the oven-loss test on 20-gram samples the index varied from 101 to 161 and in the asphaltic residue test it varied from 96 to 302. In the case of the asphaltic residue the index was low when the time of reduction was low, and for the cut-back materials the asphaltic residue test was the least severe of all tests, although there is not much difference between this and the distillation test. The index was high when the time of reduction was high, as indicated by sample 18 which took 420 minutes to come to 100 penetration and had an index of 302, and by sample 1 which took 280 minutes and had an index of 284. The index varied from 105 to 294 for 5 weeks' exposure, from 119 to 361 for 10 weeks' exposure, and from 133 to 376 for 15 weeks' exposure. Except for four samples, the index of increase was greater for 5 weeks' exposure than for any of the laboratory tests. In the exposure tests, products originally containing the highest percentages of material insoluble in naphtha generally had the smallest indexes of increase.

### OLIENSIS TEST INVESTIGATED

An interesting development in the study of asphaltic materials is the test for determining heterogeneity. This test has been called the Oliensis or spot test. The method of making the test and the interpretation of the results were outlined in a paper read before the 1933 meeting of the American Society for Testing Materials. In making this test on the original material, one part by volume of the asphaltic material was treated with 5.1 parts by volume of a special naphtha at such a temperature that solution or dispersion was complete in 6 to 8 minutes. After cooling to room temperature and adding fresh naphtha to replace any losses, a drop of the mixture was allowed to drop on a filter paper (J. H. Munktells, No. OO). The appearance of the resulting stain varied from a uniformly colored spot, in-

<sup>&</sup>lt;sup>4</sup> Report by F. J. Nellensteyn and R. Loman, Sixth Congress, Permanent Inter-lational Association of Road Congresses, first section, second question, paper 2–0.
<sup>4</sup> See Asphalts and Allied Substances by Herbert Abraham, third edition, p. 755.

dicating complete dispersion, to stains in which the center was black and rough and surrounded by a lighter-When an entirely uniform stain was obcolored ring. tained another test was made after the mixture had stood for 24 hours. The appearance of the central spot was taken as an indication of the degree of heterogeneity, and those materials that gave a uniform stain after standing 24 hours were considered as homogeneous.

The test is presumed to give an insight into the conditions of manufacture. The types of asphaltic products that should be expected to appear homogeneous

are as follows: 1. Steam-refined residuals known to have been refined without serious cracking.

2. The bitumen of certain native asphalts.

3. Some types of slightly oxidized residuals from asphaltic-base crude oils.

The materials that should be expected to appear heterogeneous are:

1. Steam-refined residuals that have been overheated during the refining process.

2. Cracking-coil residuals. 3. Highly blown residuals

The test was initially developed and used to determine whether or not petroleum asphalts had been subjected to higher temperatures than usually occur in steam refining. It has been used by some States as an identification test for the control of liquid asphaltic materials. Recently the test procedure has been standardized by a group of Middle-Western States. In applying the test to slow-curing materials they state that if the material has less than 15 percent of distillate by volume at 680° F., the test may be made on the original material. For all other materials of the slow-, medium-, or rapidcuring classes the test shall be made on the distillation The presence of the volatile distillate in these types of materials was thought to interfere with the sensitivity of the test. It will be noted that in the case of the slow-curing materials investigated the test may be run on 18 samples as received, since in only 5 cases was the percentage of distillate by volume at 680° F. more than 15 percent.

The identification of the character of the manufacturing process was the main object of this test, and for this purpose it should be made only on the finished products as they leave the refinery and not after they have been subjected to various laboratory heat tests. However, in order to determine if the character of the materials underwent a change during the laboratory tests and under exposure, the residue from the distillation test, the 20-gram oven-loss test, and the asphalticresidue test, as well as the residues from the 5-week and 15-week exposures were subjected to the Oliensis test.

For the original materials 5.1 parts of naphtha by volume were mixed with 1 part of asphaltic material. For the various residues the volume of naphtha was kept constant and the weight of a unit volume of the original material minus the weight of volatile matter that occurred in producing the residue was used. shows the ratio of naphtha to asphaltic material used for each sample. Only in the case of the asphaltic residue of sample 11 did the ratio of naphtha to asphaltic material exceed 7 to 1 by weight, the proportion that was being used by one State at the time these tests were made. It is thought that the variations in proportions of naphtha used in this work were not

TABLE 8 .- Ratio of naphtha to asphaltic material in Oliensis test

	Origi-		lation due	20-gram loss	Asphal-	Residu	
Sample identification	terial 1 (by weight)	By weight	By volume	residue (by weight)	due (by weight)	5 weeks (by weight)	15 week (by weight
	4, 10	4. 21		4. 12	5, 69	4.10	4.1
	3.74	4. 20	5, 83	4.01	5, 00	4. 13	4.2
	3.74	4.01	5, 51	3, 92	5, 22	4.00	4.1
	3.79	4.01	0.00	3.96	5, 07	3, 99	4.0
	3.96	4.58	6.04	4. 33	5. 61	4. 37	4.4
	4, 06	4.38		4, 26	6, 49	4.35	4.4
	4.11	5, 37	6, 90	4, 30	6, 37	5, 15	5.3
	4. 13	4.51		4, 27	6.32	4, 20	4.3
	3, 83	4, 29	5, 86	4. 11	5, 55	4. 16	4.5
0	4. 10	5. 28	6.84	4.92	6. 22	4.90	4.1
1	3.91	5. 22	6, 94	4. 61	7.35	4.91	5.1
2	4.10	4, 99	6, 38	4, 57	6, 68	4.75	4.
3	3, 91	4.31	5, 68	4.17	5, 86	4, 21	4.
4	3. 73	3.92		3.89	5, 36	3. 93	4.
5	4.03	4.77	6.18	4.42	6. 54	4, 44	4.
6	4. 07	4.37		4, 22	6.56	4. 26	4.
7	3, 83	4.82	6.80	4. 52	5.18	4.70	4.
8	4.13	5, 02	6, 42	4, 95	5. 52	5. 07	5.
9	3. 65	4.05	5.76	3.97	4.73	4.06	4.
20		4.78	6.05	4.53	5.95	4.65	4.
21	4 00	4.77	6.07	4. 55	6.52	4.64	4.
)*)	4.05	4, 48	5, 70	4.38	5, 76	4. 42	4.
3	3.90	4, 41	5.86	4. 29	6. 44	4. 25	4.
24	4.06	5. 40	7.40	5. 22	5. 27	5. 22	5.
25	4. 17	5. 48	7. 24	5. 38	5.34	5.36	5.
26	4. 16	5. 44	7.14	5. 36	5. 31	5. 37	5.
27		5.72	7.98	5. 54	5. 50	5.52	3.
28	4.14	5. 24	6. 90	5. 20	5. 15	5. 17	5.
29		5, 38	7, 29	5, 24	5, 24	5. 24	5.

1 All original material 5.1 naphtha to 1 of sample by volume.

Table 9.—Character of original materials and residues as deter-mined by the Oliensis test <sup>1</sup>

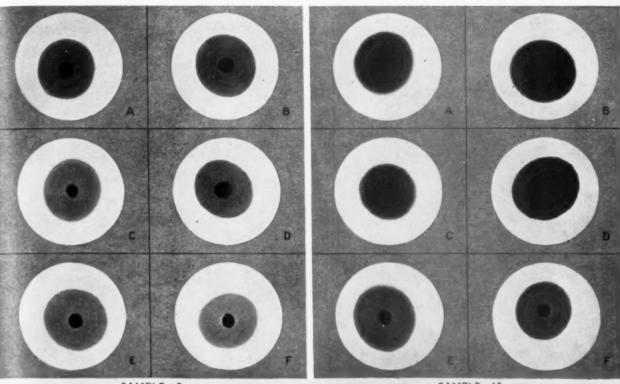
Sample identi- fication	Original material	Distilla- tion residue	20-gram loss residue	Asphaltic residue	5-week exposure residue	15-week exposure residue
	н	н	н	н	н	H
	H	H	H	H	H	11
	H	H	H	H	H	11
	H	H	H	H	H	H
	H	H	H	H	H	H
	0	0	0	0	SH	H
	SH	0	0	0	H	H
	0	SH	SH	SH	H	H
	H	H	H	H	H	H
)	0	SH	SH	0	H	H
	H	H	H	H	H	H
	0	0	SH	SH	H	H
	H	H	H	H	H	H
1	H	H	H	H	H	H
5	0	SH	SH	SH	H	H
B	SH	SH	SH	H	H	H
7	H	H	H	H	H	H
	SH	SH	0	0	SH	H
9	H	H	H	H	H	H
0	SH	SH	0	0	H	H
1	0	0	0	0	H	H
2	Ö	0	0	0	H	H
3	H	H	H	H	H	H
4	H	H	H	H	H	H
5	0	0	0	0	SH	SH
6	O	SH	SH	0	SH	SH
7	H	H	H	H	H	H
8	0	0	0	0	SH	SH
9	H	H	H	H	H	H

 $^{1}$  H = Heterogeneous; O = Homogeneous; SH = Slightly heterogeneous.

obtained. Table 9 gives a classification of the stains and figures 8, 9, and 10 show stains typical of those obtained with the various samples and their residues.

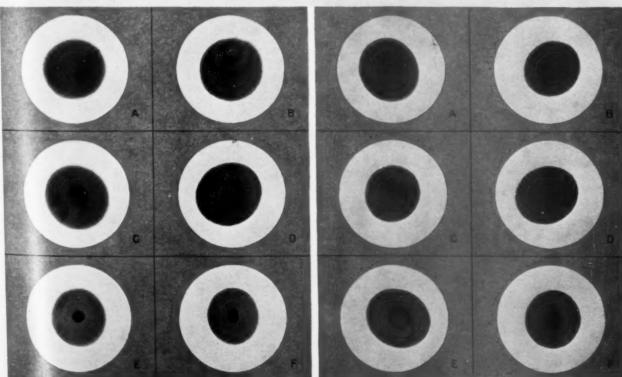
Since the results of the tests were based upon the appearance of the stain as interpreted by the observer, it is difficult if not impossible to distinguish between border-line materials or to express clearly the apparent degree of heterogeneity that may be indicated by the varying degrees of nonuniformity in the stain. classification given in table 8 should be understood to mean that, in the judgment of the observers, the materials and their residues gave stains that appeared either sufficiently wide to affect the character of the stains entirely uniform throughout or were only slightly non-

24 minimum mammam 12



SAMPLE - 2

SAMPLE - 16



SAMPLE - 8

A-ORIGINAL MATERIAL B-DISTILLATION RESIDUE

he er, en nt he he

C-20-GRAM LOSS RESIDUE D-ASPHALTIC RESIDUE

SAMPLE - 18

E-RESIDUE AFTER 5 WEEKS EXPOSURE F-RESIDUE AFTER 15 WEEKS EXPOSURE

FIGURE 8.—TYPICAL OLIENSIS STAINS.

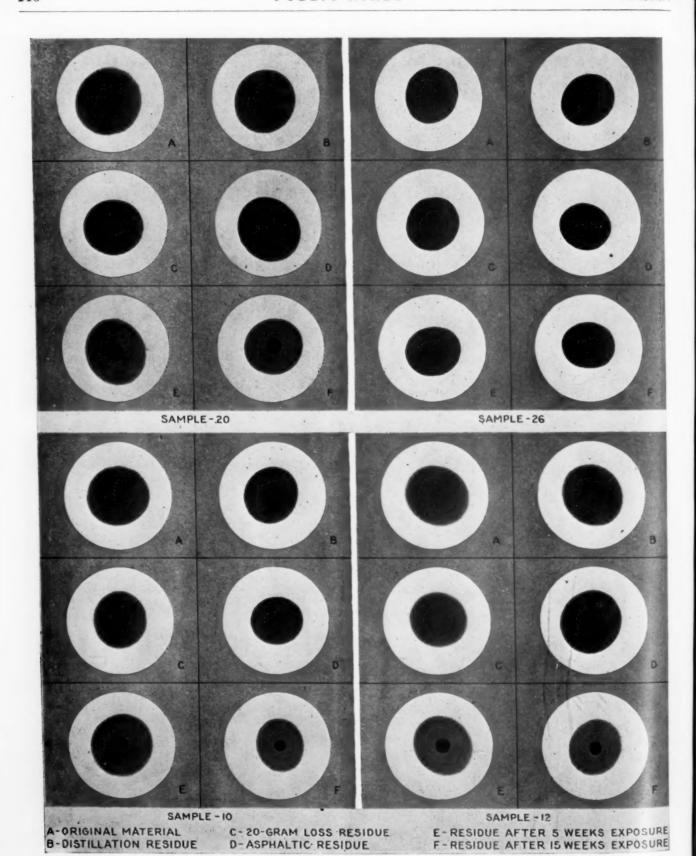


FIGURE 9.—TYPICAL OLIENSIS STAINS.

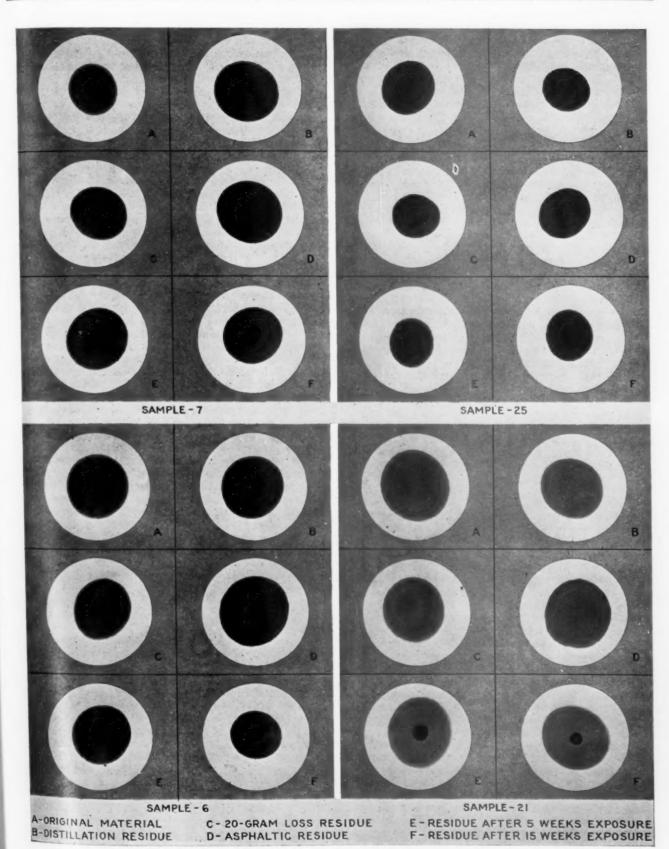


FIGURE 10.—TYPICAL OLIENSIS STAINS.

uniform, having a slightly darker, more-pronounced center, or else they had a definite dark to black center surrounded by a uniformly lighter-colored stain and were consequently classified as homogeneous, slightly heterogeneous, and heterogeneous, respectively. No attempt was made to indicate the extent or degree of heterogeneity other than to classify as slightly heterogeneous those materials and residues giving stains only slightly nonuniform.

### ALL EXPOSURE RESIDUES FOUND TO BE HETEROGENEOUS

A study of the results of the Oliensis test shows that of the 29 materials, 10 were homogeneous, 4 slightly heterogeneous, and 15 definitely heterogeneous in their original state. All the residues from exposure were heterogeneous or slightly heterogeneous. Of the 10 materials appearing homogeneous in their original state, 5 developed homogeneous residues in all 3 laboratory loss tests, 2 developed slightly heterogeneous residues in all 3 tests, 2 developed slightly heterogeneous residues in the oven test and distillation test but not in the asphaltic-residue test, and 1 developed slightly heterogeneous residues in the oven test, and asphaltic-residue tests but not in the distillation test.

Of the 4 materials appearing slightly heterogeneous in their original state, 1 developed homogeneous residues in all 3 laboratory tests, 2 developed homogeneous residues in the asphaltic-residue and loss tests but remained heterogeneous in the distillation test, and one material remained heterogeneous in all tests. The 15 materials that appeared heterogeneous in their original state all developed heterogeneous residues.

A comparison of these materials according to the character of the stains produced in the laboratory and their behavior under laboratory and exposure conditions may be of interest. Nine of the materials, samples 2, 3, 4, 9, 14, 17, 19, 23, and 27, were heterogeneous originally and had heterogeneous residues. The photograph of sample 2 in figure 8 is typical. All of the above samples, except 23 and 27, showed micropscopic flecks as shown in figure 2 when examined under the microscope. All except samples 14 23, and 27 had appreciable amounts of material insoluble in carbon tetrachloride in the original sample. Samples 14, 23, and 27 developed carbenes at the end of 15 weeks of exposure and the other samples showed an increase in carbenes and free carbon.

All of the nine samples, except sample 23, had high percentages of material insoluble in naphtha. All had specific gravities greater than 1.01, except sample 27. This material has a very high specific gravity for a fluid cut-back and its behavior and characteristics place it in this group having high specific gravities. The asphaltic residues of the 9 samples showed the effect of changes in temperature, having low penetration at 32° F. and no ductility at 34°-35° F., although all had ductilities of over 110 at 77° F.

After 15 weeks' exposure all the materials were very hard, except sample 9, which, at the end of 5 weeks, had separated into two parts, one hard and brittle, the other It was impossible to flux these two parts soft and oily. so that, while having a float at 122° F. of over 1,000 seconds at the end of 5 weeks, it was impossible to get a penetration until the end of 15 weeks. When running the softening point test on the residues of this sample, after exposure, the material did not flow slowly to the bottom but dropped immediately at the temperature

sample but the results are unimportant as no true ductility value was obtained. The residues after 15 weeks' exposure of samples 2, 3, 4, 17, 23, and 27 had very low ductility at 77° F. but samples 14 and 19 had ductilities at 77° F. of 97 and 110+, respectively. None of the residues had any ductility at 34°-35° F.

Three of the materials, 5, 11, and 13, gave Oliensis stains identical with those of the preceding samples. They did not, however, have as high specific gravities. were clear in the microscopic test, and did not, even at the end of 15 weeks, develop any carbenes. asphaltic residues had some ductility at 34°-35° F. and, at the end of 15 weeks, while having low ductility at 77° F., samples 5 and 13 had some ductility at 34°-35° F

The remaining 17 materials all had low specific gravities, were clear in the microscopic test, did not have carbenes, and, with 1 exception, did not develop them. They showed various types of stains in the Oliensis test although all were heterogeneous at the end of 5 Their asphaltic residues varied in ductility and effect produced by changes in temperature as did their residues after exposure.

The stains of samples 1, 24, and 29, while resulting in photographs similar to those of the high-gravity materials, did not have the nuclei as raised or as rough as the stains of the materials with high specific gravities. The producers of the cut-back asphalt designated as sample 24 stated that their plant had no cracking equipment. This material, therefore, had evidently become heterogeneous in the refining process because of overheating, since the residue obtained in laboratory tests had good ductility, indicating that it had not been overblown. Sample 29 likewise was a cut-back asphalt with a ductile base. Sample 16 was slightly heterogeneous originally, produced slightly heterogeneous residues by distillation and volatilization, and produced a heterogeneous asphaltic residue. The asphaltic residues of these four samples were not especially affected by temperature change, having some ductility at 34°-35° F., although the asphaltic residues of samples 1 and 16 had comparatively low ductilities at 77° F. The residues of samples 1 and 16 after exposure had low ductility at 77° F. Those of samples 24 and 29 (cutback asphalts) were very hard and consequently had very low ductility.

### OLIENSIS TEST MORE SENSITIVE THAN MICROSCOPIC TEST IN DETECTING OVERHEATED MATERIALS

Samples 8 and 15 were homogeneous but all of their residues from laboratory tests were slightly heterogeneous. The asphaltic residue of sample 8 had relatively low ductility at 77° F. but that of sample 15 had good ductility. Both asphaltic residues had good ductilities at 34°-35° F. The residue of sample 15 after exposure had low ductility at 77° F. Sample 8, the only material of low specific gravity to develop carbenes, separted in the exposure test in the same manner as sample 9.

Samples 18 and 20 were both slightly heterogeneous but their asphaltic and loss-test residues were homogeneous. Their asphaltic residues and their residues after exposure had good ductility at 77° F. Although the asphaltic residue of sample 20 had a ductility of only three-fourths centimeter at 34°-35° F., the ductility of its residue after exposure, as well as the ductility of the two residues of sample 18, was good at 34°-35° F. reported in table 5. Ductility tests were made on this Sample 18 developed a residue at the end of 5 weeks'

man of 111 1111 to 100

exposure that was the least heterogeneous of any of the residues from the slow-curing materials.

Samples 10 and 26 and their asphaltic residues were homogeneous. The asphaltic residues of both samples had good ductility at 77° F. and 34°-35° F. At the end of 15 weeks, sample 10 had low ductility at 77° F. but good ductility at 34°-35° F., while sample 26, a cut-back asphalt, was extremely hard and nonductile.

Sample 12 was homogeneous as was its residue after distillation. Its asphaltic residue had good ductility at 77° F. and 34°–35° F. and the residue after exposure had fair ductility at 77° F. and good ductility at 24° 25° F.

Sample 7 was slightly heterogeneous but all of its residues from laboratory tests were homogeneous. Its asphaltic residue was ductile at 77° F. but only slightly so at 34°-35° F. and its residue after 15 weeks of exposure had good ductility at 77° F. and at 34°-35° F. Samples 6, 21, 22, 25, and 28 were homogeneous with

Samples 6, 21, 22, 25, and 28 were homogeneous with homogeneous residues from laboratory tests. All of their asphaltic residues had good ductility at 77° F. and all except those of samples 21 and 22, the California residuals, had good ductility at 34°–35° F. At the end of 15 weeks, sample 6 was still fluid, while the cutback asphalt samples 25 and 28 were very hard and nonductile. Samples 21 and 22 had good ductility at 77° F. and at 34°–35° F. although their asphaltic residues were nonductile at 34°–35° F.

It is readily apparent that the laboratory tests did not produce residues that gave stains in the Oliensis test radically different from the stains of the original ma-The behavior of the residues from exposure showed, as did the other tests, that outdoor exposure alters asphaltic materials far more than any of the laboratory heat tests. This was strikingly shown by the decidedly heterogeneous stains obtained with the residues from exposure, especially in the case of the materials originally homogeneous. It is not believed, however, that it is possible to predict the physical and chemical characteristics of the material after exposure from the results of the Oliensis test, whether made on the original material, the residues from laboratory tests or both. Residues having what are believed to be desirable qualities were obtained from both homogeneous and heterogeneous materials, although heterogeneous materials undoubtedly have a more pronounced tendency to carbonize and their slow-curing products generally develop a less-ductile residue.

For detection of materials that have been inadvertently or intentionally subjected to too high a temperature during the refining process, the Oliensis test seems to be more sensitive than the microscopic test. All of the materials that had the characteristics of overheated or cracked materials were heterogeneous in the Oliensis test but only seven of them showed microscopic flecks.

### HUBBARD-FIELD STABILITY TEST USED TO MEASURE BONDING STRENGTH AND DEVELOPMENT OF BONDING STRENGTH UPON EXPOSURE

Cylinders were made according to the Hubbard-Field method and tested to determine the adhesiveness or bonding strength of the original material, the residue after distillation and the asphaltic residue, and the development of bonding strength by the original materials after exposure. The first series, for the determination of bonding strength, consisted of 3 sets of 3 cylinders each for each material. The first and second

sets contained 16.6 percent by volume of the original material and distillation residues respectively mixed with 83.4 percent of a standard sand. The third set contained the same percentage of asphaltic residue by weight as was contained in the cylinders made with the original materials that gave an almost constant percentage of bitumen by volume in the cylinders of this set. All cylinders of series 1 were tested immediately for stability at 77° F.

The second series of cylinders, for determination of the development of bonding strength, likewise consisted of 3 sets of 3 cylinders using the same aggregate used in the first series and the same percentage of the original materials by volume. These three sets were placed in the exposure boxes and subjected to the same exposure conditions as the thin films. One set was removed at the end of 5, 10, and 15 weeks. The cylinders were weighed before and after exposure and the loss in weight was expressed as a percentage of the bituminous material present in the cylinder as made. After weighing, the cylinders were tested for stability at 77° F.

For comparative purposes two additional sets of cylinders were made, using as a binder the amounts of distillation residue and asphaltic residue that would have been obtained if the bitumen in the cylinders containing the original material had been subjected to the distillation or asphaltic-residue test. The aggregate used was a Potomac River sand that had been separated on standard sieves and recombined to give the following grading:

P	erces	nt
Passing no. 10, retained on no. 20	3.	7
Passing no. 20, retained on no. 30	10.	3
Passing no. 30, retained on no. 40	18.	1
Passing no. 40, retained on no. 50	21.	3
Passing no. 50, retained on no. 80	36.	6
Passing no. 80, retained on no. 100	6.	1
Passing no. 100, retained on no. 200	3.	2
Passing no 200		7

This sand had a specific gravity of 2.666 and the voids in the mineral aggregate, determined on the compacted cylinders of both series, were 38 percent for the cylinders made with the original materials, 37.4 percent for the cylinders made with the distillation residue, and 36.9 percent for the cylinders made with the asphaltic residue.

The method of mixing and molding the cylinders was the same as that used in 1932. The results of the tests on the cylinders of series 1 and 2 are given in tables 10 and 11, respectively. All results are the averages of three tests.

The results of tests on the cylinders of series 1 are shown graphically in figure 11. The stability of the cylinders at 77° F. was plotted against the Furol viscosity at 122° F., and the results of the float test at 77° F. for the cylinders made with the original materials and against the float test results at 122° F. and the penetration at 77° F. for the cylinders made with the distillation residue. Since the asphaltic residues are all of approximately the same consistency, the stabilities were plotted for each sample independently.

Figure 11 shows that although the stability of the mixtures was roughly proportional to the consistency of the contained bitumen, materials having the same consistency as measured by viscosity at 122° F., float test at 77° F. and 122° F., and penetration at 77° F. had different stabilities. This was especially noticeable

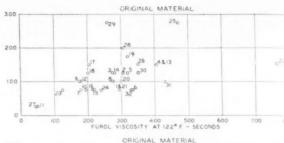
Table 10.—Results of tests on series 1 cylinders

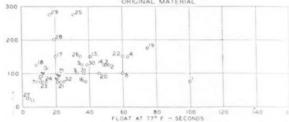
	Orig	inal mate	erial	Distil	lation re	sidue	Asphalti	e residue
Sample iden- tification	Stabil- ity at 77° F.	Float at 77° F.	Furol viscos- ity at 122° F.	Stabil- ity at 77° F.	Float at 122° F.	Pene- tration at 77° F.	Stabil- ity at 77° F.	Pene- tration at 77° F
				-	-	-		-
	Pounds	Seconds	Seconds	Pounds	Seconds		Pounds	
	75	100	294	125	24		2, 275	9
	125	52	303	400	60		3, 975	10
	125	50	274	200	43		3, 775	10
	150	63	404					
				250	45		3,600	10
	125	36	331	325	58		2, 575	10
	75	21	335	150	21		2, 275	9
	75	9	181	375	62		2, 575	9
	100	60	277	100	27		2, 100	16
)	100	33	175	200	44		3,500	8
0	75	11	197	475	87		2, 425	10
1	25	4	50	175	34		3, 175	8
2	100	13	182	200	38	*******	2, 450	9
3	150	41	406	200	37	*****	2, 775	9
14	125	49	278	150	36			
4	75					*****		
15		21	219	175	41		2, 675	10
16	75	38	217	100	25		2, 225	10
17	150	20	205	1,000	110		2,700	10
18	125	9	199	400	60		1,900	16
19	175	75	320	475	57		3,775	10
20	100	46	300	350	50		2, 325	16
21	75	23	293	200	41		2, 375	1
22	150	60	768	325	56		1 0 000	1
23	75	12	123	150	27		2,700	1
24		14	241	3, 925		76	2, 950	1
25	275	31	472	3, 225		. 77	2, 825	1
26			351	3,025	******	. 84	2,725	
27			47	3,800			3,850	
28	. 200		303	3,775	lanners.	65		
29	275	17	260	3,850		. 72	3, 650	1
30	125					. 81		î
31	100		431	2, 325		205		î
32	-					298		1

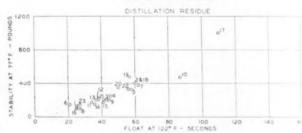
TABLE 11 .- Results of tests on series 2 stability cylinders

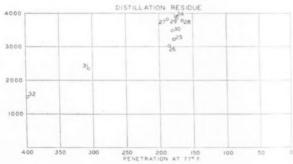
	Cylin	nders ma	de with	the or	iginal :	materi	als	Cylin made distill resid	with ation	Cylin made aspha resid	with
	St	tability s	t 77° F.		Loss	of bitu	men	Sta-	Theo- reti-	Sta-	Theo-
outinities and management	When made			In 15 weeks		In 10 weeks		bility at 77° F.	cal loss of bitu- men	bility	cal loss of bitu- men
					Per-	Per-	Per-		Per-		Per-
	Lbs.	Lbs.	Lbs.	Lbs.	cent	cent	cent	Lbs.	cent	Lbs.	cent
2	75	300	300	425	0	3	1	125	3	1, 975	25
3	125 125	900 750	1, 100	1,300	7	12	12	425	11	3, 575	25
4	150	600	1, 000 750	1, 150 850	6	10	11	250	7	3, 075	25
5	125	675	1, 025		3	6	6	300	6	3, 150	2
6	75	225	250	1, 275	6	11	11	475 1 175 425 2 175 325 1 550 2 275 2 275 1 275 225	14	2, 400	2
7	75	425	550	300 600	15	9	9	175 425 275 325 1550 275 275 275 275 225	7	1,975	38
8	100	250	300	300	3	20	21	1 425 2 5 175 2 2 325 1 2 550 2 2 275 2 5 275 1 0 275 2 3 300 1 9 175	23	2,075	3.
9	100	550	775	800	7	12		6 175 2 325 1 550 2 275 2 275 1 0 275 8 225 3 300 9 175	6	1, 750	3.
)	75	775	950	1, 250	14			6 175 2 325 1 2 2 325 2 2 275 2 2 275 3 2 2 2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3		3, 075	3
	25	425	500	800	18	18 21		12		2, 175	3
	100	350	425	550	10			2 325 11 8 550 22 275 24 15 275 16 8 225 13 300 15 9 175		2, 450	4
	150	650	700	875	6	14		18 550 22 22 275 24 15 275 16 10 275 16 8 225 13 300 16		2, 050	3
	125	650	900	1, 025	4	8		22 275 21 15 275 18 10 275 6 8 225 1 13 300 1 9 175	9	2, 350	3
,	75	475	550	600	8	12			5	2,900	3
3	75	250	325	350	4	8				2, 325	3
7	150	950	1,000	1, 200	15	19				1,875	3
3	125	825	950	1, 200	16	20				2, 325	2
9	175	1,650	2, 525	3, 075	7	11		21 1, 275 21 20 500 18 11 625 16 13 350 14		1,750	1 3
0	100	725	1,000	1, 250	8	12		20 500 18 11 625 10 13 350 14 14 300 18		2, 825	1
1	75		700	850	8	13		11   625   10 13   350   14 14   300   13 9   350   10 11   200   13		2, 050 1, 950	3
2	150	775	1,075	1,400	5	9		14 300 18 9 350 10 11 200 12		3	
3	75	275	550	750	7	11		9 350 10 11 200 13 22 3,650 2		2, 100 2, 175	1 2
14	75		4, 325	5, 200	20	23	22	11 200 12 22 3,650 29 19 3,025 2			3
25			3, 750	3, 850	17	19	19	2 3, 650 2 9 3, 025 2 3 2, 800 2		2,800	1 3
26	150	3, 500	3, 125	3, 550	21	24	23	3, 650 2 3, 025 2 3 2, 800 2 2, 875 3	23	2,650 2,775	
7		1,450	1,500	1,650	28	30	29	3, 025 2 2, 800 2 2, 875 3 3, 400 2	30	3, 600	1 :
8	200	4, 650	4, 250	4, 975	16	18	18	3, 025 2 2, 800 2 2, 875 3 3, 400 2	21		
9		4, 400	4, 100	4, 825	13	16	16	3, 025 2 2, 800 2 2, 875 3 3, 400 2	24	3, 475	1
0				-,		-0	20	3, 175	26		
1								2, 075	20		1
2	75		1					1 495	20		

in the results with the cylinders made with the asphaltic residue. Although all of these residues had approximately the same penetration, the stability of the cylinders varied from 1,900 pounds for sample 18 to 3,975 pounds for sample 2.









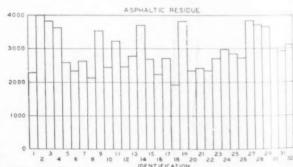


FIGURE 11.—RELATION BETWEEN THE CONSISTENCIES OF ORIGINAL MATERIALS, DISTILLATION RESIDUES, AND ASPHALTIC RESIDUES AND THE STABILITY AT 77° F. OF CYLINDERS OF SERIES 1.

It is seen that the cylinders made with the asphaltic residues of samples 2, 3, 4, 9, 11, 14, and 19 all had stabilities of over 3,000 pounds. All of these materials were heterogeneous originally, all were materials of high specific gravity, and all except sample 11 contained or developed carbenes and free carbon. Sample 11 had a relatively high specific gravity but did not develop

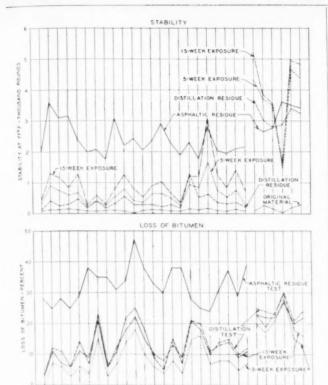


FIGURE 12.—Comparison of Loss of Bitumen and Stability of Series 2 Hubbard-Field Cylinders.

carbenes. Samples 5, 7, 13, 15, 17, and 23 had stabilities between 2,500 and 3,000 pounds. Samples 17 and 23 were heterogeneous materials of high specific gravity that contained or developed carbenes. Samples 5 and 13 were heterogeneous materials of fairly high specific gravity but they did not develop carbenes, and samples 7 and 15 were materials of low specific gravity. All of the cut-back products had asphaltic residues giving stabilities of 2,500 pounds or over and five gave stabilities of 3,000 pounds or over.

Figure 12 shows the results of stability tests on the cylinders of series 2. The loss of bitumen in 5 and 15 weeks of exposure and the theoretical loss of bitumen in the cylinders made with the distillation and asphaltic residues were plotted for each sample. The stabilities at 77° F. for each sample were also plotted.

In this figure it is seen that, in the case of the slow-curing materials, although the loss of bitumen in the exposed cylinders was approximately the same as the loss in the distillation test, the exposed cylinders had greater stability than the cylinders made with the distillation residue except in the case of sample 17. This sample, even in 15 weeks, did not attain as high a stability as the cylinders made with the distillation residue. It is also seen that the loss in 15 weeks' exposure did not approach the loss in the asphaltic residue test and that the stability of the exposed cylinders did not approach the stability of the cylinders made with the asphaltic residue, except in the case of sample 19.

For the cut-back materials the indicated losses were probably in error due to unavoidable loss of volatile matter while mixing and molding the cylinders. The losses in 5 and 15 weeks of exposure probably should have been about the same as the losses in the distillation and asphaltic residue tests. The stabilities at 5 weeks

were higher than the stabilities of cylinders made with the asphaltic and distillation residues except in the case of sample 27.

### SATISFACTORY CHECKS OBTAINED WITH RESULTS OF 1932 TESTS

After the exposure tests had been started, a question was raised concerning the use of plate glass covers for the exposure cabinets because it prevented the active ultra-violet rays from acting on the materials. Fused quartz glass not being available, Vita glass, which, after a short stabilization period, is guaranteed to permanently transmit an effective volume and combination of wave lengths of active ultra-violet light, was used to determine the effect of the passage of more active light. Duplicate sets of 10 of the slow-curing materials, 2 of the rapid-curing materials, and sample 27 and 3 new medium-curing materials, samples 30, 31, and 32, were exposed under both Vita and plate glass for 5- and 10-week periods. The materials were exposed in thin films and also admixed with the standard sand in the form of Hubbard-Field cylinders.

This exposure was started August 28, 1933. During the first 5-week period the average maximum air temperature was 80° F. and the number of sunlight hours was 266. During the 10-week period the average maximum air temperature was 73° F. and the number of sunlight hours was 512. The results of the tests on the thin films are given in table 12, and those on the Hubbard-Field cylinders in table 13.

As shown by tables 6 and 12, the materials did not lose as much nor get as hard in 10 weeks as they did in the original 5 weeks of exposure. This was due to the lower air temperature and also to the fact that the sun's rays striking at an oblique angle did not cause the material to get as hot as earlier in the summer. There was little if any difference between the materials exposed under the different types of glass. The samples exposed under Vita glass had a little more material insoluble in carbon disulphide and carbon tetrachloride and generally had a little more material insoluble in naphtha. Each solubility reported was the average of 3 or more tests. The results of the stability tests do not show that there was any difference between the two types The comparative study of the effectiveness of Vita glass and plate glass as cover for the exposure of the samples did not produce differences great enough to indicate their relative efficiency for this purpose.

As stated previously, four of the samples tested in 1932 were included in the 1933 work and the results obtained, as shown in table 14, were in remarkably close agreement with the previous tests. In the case of samples 17 and 19 the residues from the 1933 exposure tests had greater percentages of free carbon and carbenes than did residues from the 1932 exposure tests.

The results of the two sets of stablity tests were not in such close agreement because the aggregate used in the 1933 tests was somewhat coarser than that used in 1932. In 1932, cylinders made with sample 17 were the only ones that, after 15 weeks' exposure, had a stability about the same as those made with the distillation residue. In 1933 the cylinders made with sample 17, after 15 weeks' exposure, had less stability than the cylinders made with the distillation residue. Cylinders of sample 19 that in 1932, after 15 weeks' exposure, had a stability approaching that of the asphaltic-residue cylinders, in 1933, after 15 weeks' exposure, had a stability higher than that of the asphaltic-residue specimens.

Table 12.—Results of tests on plate and Vita glass exposure residues

### 5 WEEKS' EXPOSURE

-					Plate	glass									Vita	glass				
ntification		2	Peneti	ration	point		ity at 5 minute	atter in- n CS s	tter in-	186° В.		Sa <sub>4</sub>	Penetr	ation	point		ity at 5 minute	20	matter in-	St. R
Sample identification	Loss	Float at 122°	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening p	At 77° F.	At 34°- 35° F.	Organic mat soluble in	Organic matter in- soluble in CCI+	Insoluble in 86° naphtha	Loss	Float at 122°	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening p	At 77° F.	At 34°- 35 ° F.	Organic mat soluble in	Organic ma soluble in	tasoluble in se
	Per- cent -2.5	Sec- onds 50			°F.	Centi- meters		Per- cent	Per- cent	Per- cent 12.7	Per- cent -3.3	Sec- onds 46			°F.	Centi- meters	Centi- meters	Per- cent	Per- cent	P
3	6.7 4.5	113 61						0.82	6.86	29. 9 18. 4	4.5	92 54					*****	1. 22	6.85	
1 7 8	2. 2 12. 4 16. 0	53 140 110						1.08	2. 68 4. 09	24. 8 26. 8 12. 2	1. 6 12. 4 14. 9	47 156 104						1. 28	2.73 4.64	
9 0	6.4	105 103						1.68	9. 63	32.6 17.2	5. 6 6. 5	98 96						2.00	10. 22	
3	8.0 6.0 21.5	53 52	37	18	143	8. 5	0.0	. 13	. 33	14, 1 19, 2 27, 3	5.7 5.6 21.5	49 50	33	16	147	10	0.0	. 26	. 15	
9	27. 4 18. 5	******	46	14	124 141	17. 0 11. 0	.0	. 19	. 56	27. 7 33. 3 32. 7	26. 2 18. 8 22. 1		39 59 40	14 21 20	126 134 138	12 18 15	.0	. 31	. 64	
0 1 2	22. 6 17. 6 17. 7		38 61 122	16 18 24	136 124 106	18.0 110+ 110+	.0			29. 7 19. 8	17. 3 17. 5		52 123	17 28	126 106	110+ 110+	.0			
								10 WE	EEKS' I	EXPOS	SURE									
	-2.5 7.1	54 152						1. 42	0. 14 7. 77	14. 1 30. 2	-3.3 4.1	56 141		******				1.73	0. 22 8. 13	
3		85							. 04	19.6	3. 2	71							. 05	
4	2.2	64						. 50	3. 16	23. 9	2.3	65						. 55	3.45	
7	12.9	178						1.03	5. 16	28. 1	12.6	297						1.24	5. 39	
8		144						0.00	. 12	12.9	16. 1	154						9.00	. 16	
9	7.6	145						2.92	10. 42	34. 3 19. 4	6. 9 8. 0	133 125						2.99	10.61	
01	8.3	135							.07	15. 0	8.7	74	******	*****					.11	
3	6.1	71							. 10	20. 4	6.1	84	******	******					.08	
6		41	32	17	147	6.3	.0	. 42	. 36	28. 0	21.5	0.8	30	16	149	6	.0	. 31	, 25	
7	26. 4		0.1		128	8.5		. 25	. 73	29. 9	26.7		26	13	130	7	.0	. 30	. 59	
9	19. 3		42	17	141	13. 5			. 22	33. 4	18.9		52	19	136	16	.0		. 24	
			1 20	0.4									34							
0	_ 22.6		32	14	142	8.3	.0		. 43	33.9	22.3		34	15	141	11	.0		. 20	
0 1 2			1 00	14	142 131 115	33. 0 63. 0	.0		. 13	33. 9 33. 7 22. 2	17. 3 18. 0		44 73	15 15 17	131	32 65	.0	******	.14	

### CONCLUSIONS

The results of this investigation substantiate most of the conclusions arrived at in the 1932 investigation, modify two of the conclusions, and indicate some new conclusions. The conclusions substantiated are:

1. Materials of high specific gravity and their residues are, in general, more susceptible to changes in temperature than materials of low specific gravity and their residues.

2. Hardening due to causes other than loss of volatile matter, and changes in inherent characteristics that may be attributed to oxidation, polymerization, and carbonization, occur to the greatest extent upon exposure and least during distillation.

3. The development of a ductile residue either in the asphaltic-residue test or, in the case of cut-back materials, in the distillation test, does not indicate that the material will develop a ductile residue upon exposure.

4. The bonding strength of the original materials and their residues is roughly proportional to their consistencies, but materials having the same consistency as measured by the present tests do not always give the same stability. The reasons for these differences in stability cannot be determined under the present methods of testing.

The following conclusions are somewhat modified from 1932:

Table 13.—Results of stability tests on cylinders exposed under plate and Vita glass

		Loss of	bitumen		1	Stability	at 77° F.	
Sample identi- fication	In 5	weeks	In 10	weeks	In 5 v	weeks	In 10	weeks
	Plate	Vita	Plate	Vita	Plate	Vita	Plate	Vita
	Percent	Percent	Percent	Percent	Pounds	Pounds	Pounds	Pound
1	2	1	2	1	225	250	275	37
2	8	5	8	7	850	700	850	80
13	7	4	6	5	450	450	500	42
14	4	2	4	3	475	400	500	45
17	13	12	15	13	650	700	650	70
18	18	16	18	17	500	500	525	50
19	8	6	8	8	1.075	1,000	1,250	1, 27
20	8 8	7	8	8	550	500	500	52
21	8	7	8	8	325	350	350	35
23	5	4	6	5	225	200	250	25
26	21	21	24	23	2,775	2,650	3,550	3, 50
27	28	28	30	30	1, 225	1, 225	1,575	1, 35
29	12	12	14	13	3, 775	3, 725	4,600	4, 57
30	23	23	24	24	2, 575	2,550	2,850	3, 02
31	18	19	20	20	2, 325	2,375	2,725	3, 12
32	18	17	19	20	1.675	1,725	2, 200	2,67

5. The relative rates of volatilization of the various materials can be anticipated most readily from the distillation curves. The different classes of material may be differentiated in the loss and asphaltic-residue tests, especially if the time of reduction to 100 penetration is considered. However, sharp distinctions in initial curing properties, that may be of importance in some types of construction, can be determined only from the distillation curves.

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6. Carbonization generally occurs in materials that originally contain some material insoluble in carbon disulphide and carbon tetrachloride, but some materials with exceptionally high solubility in these solvents show a tendency to carbonize under both laboratory and exposure conditions.

The following conclusions are developed upon the basis of the data collected in 1933 only:

7. The Oliensis test is more sensitive than the microscopic test in the detection of materials that have been subjected to excessively high temperatures in manufacture. However, neither test seems definitely to distinguish products that will weather badly.

8. The use of Vita glass in place of plate glass for the cover of the exposure boxes did not materially change the results. However, because of the lateness in the year when these tests were started the results are considered inconclusive.

9. If like periods of the year are used for exposure, satisfactory check tests can be obtained with the exposure assembly used in these investigations.

Many of the laboratory heat tests have been criticized as producing conditions dissimilar to and more severe than service conditions. These investigations have shown that the physical and chemical characteristics generally believed to belong to unsatisfactory materials are developed upon exposure in many products that satisfactorily withstand laboratory testing. While it is possible, by the utilization of identification tests, to restrict materials to a limited number of sources or manufacturing processes, it is impossible to predict, with any degree of accuracy, the weather-resisting properties of the material thus obtained. It is believed that efforts should be directed to the modification of some of the present laboratory heat tests so that differences in the tendency of various materials to develop unsatisfactory residues may be recognized.

Table 14.—Comparison of 1932 and 1933 exposure tests
TESTS ON RESIDUES AFTER EXPOSURE

			Sam	ple ide	ntifica	tion		
Test characteristic	1	7	1	8	15	9	2	0
	1932	1933	1932	1933	1932	1933	1932	1933
Loss in 10 weeks, percent Penetration at 77° F Penetration at 32° F	18.3	18.7	18. 5	17. 7	13.3	12.0	13. 1	11.5
Penetration at 77° F	41	36	132	152	40	44	95	136
Penetration at 32° F	17	14	37	43	9	10	30	41
Softening point, F.	163	170	106	105	121	121	115	115
Ductility at 77° F., centimeters. Ductility at 34°-35° F., centi-	1.0	0.5	110+	110+	110+	110+	65. 0	90.0
meters	. 0	.0	1.5	7.5	.0	.0	3.3	3.8
Insoluble in C82, percent	1.06	3, 17			5, 96	6, 50		
Insoluble in CCls, percent	8. 27	9.66	. 10		14.94	16, 02	. 05	
Insoluble in 86° B. naphtha,								
percent	34. 8	36. 2	17.6	17.3	41.1	38.7	26.8	24.
Loss in 15 weeks, percent	19, 8	19.5	18.7	19.0	13.7	13.8	13.7	13.
Penetration at 77° F	19	29	93	93	28	29	67	8
Penetration at 32° F	12	12	29	28	8	7	24	2
Softening point, °F Ductility at 77° F., centi-	181	168	113	116	129	128	121	12
Ductility at 77° F., centi-								
meters	0.3	0.5	86.0	115.0	110+	110+	42.0	72.
Ductility at 34°-35° F., centi-								-
meters	.0	.0	4.0	4.5	.0	. 0	1.3	3.
Insoluble in CS2, percent	2, 50	3.30			6. 17	7. 50		
Insoluble in CCl4, percent	9.11	9.88	. 17	. 10	15.61	18. 15	. 14	. 0
Insoluble in 86° B. naphtha,								
percent	37.2	35. 4	19.6	18.8	38.8	39.9	28. 5	27.

TESTS ON HUBBARD-FIELD CYLINDERS

Loss of bitumen:								
In 5 weeks, percent	15	1.5	18	16	8	7	9	8
In 10 weeks, percent	18	19	19	20	10	11	12	12
In 15 weeks, percent	21	21	21	20	14	11	13	13
Stability at 77° F.:1								
Original cylinders, pounds.	275	150	200	125	325	175	225	100
After 5 weeks' exposure,								
pounds	1, 100	950	800	825	1, 375	1,650	775	725
After 10 weeks' exposure.						1		
pounds	1,575	1,000	1, 125	950	3, 175	2, 525	1, 525	1,000
After 15 weeks' exposure.								
pounds	1.550	1, 200	1,650	1, 200	4, 050	3,075	1.975	1, 250
Series 2, distillation residue	-,			-	1			
cylinders, pounds	1.475	1, 275	750	500	800	625	500	350
Series 2, asphaltic residue				1	1			
cylinders, pounds	4, 050	2 325	2.925	1.750	4, 900	2,825	3, 325	2, 050

Differences in stability probably are caused by differences in grading of the sand.

Table 12.—Results of tests on plate and Vita glass exposure residues

### 5 WEEKS' EXPOSURE

					Plate	glass									Vita	glass				
ntification		F.	Penetr	ration	point	Ductili cm per	ty at 5 minute	tter in- CS 2	tter in-	86° B.		E4 .	Penet	ration	oint		ity at 5 minute	30	ccl.	NG. 13.
Sample identification	Loss	Float at 122°	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening p	At 77° F.	At 34°- 35° F.	Organic matt soluble in	Organic matter in soluble in CC14	Insoluble in 86° naphtha	Loss	Float at 122° F	At 77° F., 100 g, 5 sec.	At 32° F., 200 g, 60 sec.	Softening point	At 77° F.	At 34°- 35 °F.	Organic mat soluble in	Organic matter in soluble in CCI 4	Insoluble in
	Per- cent -2.5	Sec- onds			°F.	Centi- meters	Centi- meters	Per- cent	Per- cent	Per- cent 12.7	Per- cent -3.3	Sec- onds 46			°F.	Centi- meters	Centi- meters	Per- cent	Per- cent	F
************	6.7	113				******		0.82	6.86	29.9	4.5	92	******					1. 22	6.85	
· · · · · · · · · · · · · · · · · · ·	4. 5 2. 2 12. 4	61 53 140	*******					. 29 1. 08	2.68 4.09	18. 4 24. 8 26. 8	2. 2 1. 6 12. 4	54 47 156			******		******	.47 1.28	2.73 4.64	
	16.0 6.4 7.9	110 105 103						1.68	9. 63	12. 2 32. 6 17. 2	14.9 5.6 6.5	104 98 96						2.00	10, 22	
	8.0 6.0 21.5	53 52	37	18	143	8.5	0.0	. 13	. 33	14. 1 19. 2 27. 3	5. 7 5. 6 21. 5	49 50	33	16	147	10	0.0	. 26	. 15	
	27. 4 18. 5 22. 6 17. 6 17. 7		46 40	14 17 16 18 24	124 141 136 124 106	17. 0 11. 0 18. 0 110+ 110+	.0	. 19	. 56	27. 7 33. 3 32. 7 29. 7 19. 8	26. 2 18. 8 22, 1 17. 3 17. 5		39 59 40 52 123	14 21 20 17 28	126 134 138 126 106	12 18 15 110+ 110+	.0	.31	. 64	
								10 W F	EKS'	EXPOS	URE									
*************	-2.5 7.1 5.1	54 152 85						1. 42	0.14 7.77 .04	14.1 30.2 19.6	-3.3 4.1 3.2	56 141 71						1. 73	0. 22 8. 13 . 05	
************	12.9 16.4	64 178 144 145						1. 03 2. 92	3. 16 5. 16 . 12 10. 42	23. 9 28. 1 12. 9 34. 3	2.3 12.6 16.1 6.9	65 297 154 133						2. 99	3. 45 5. 39 . 16	
	. 8.3 8.7	135 70 71							.07	19. 4 15. 0 20. 4	8. 0 8. 7 6. 1	125 74 84						2. 99	10. 61 , 11 , 12 , 08	
***************************************	21. 4		32 31 42	13	147 128 141	6. 3 8. 5 13. 5	.0	. 42	. 36	28. 0 29. 9 33. 4	21. 5 26. 7 18. 9		30 26 52	16 13 19	149 130 136	6 7 16	.0	.31	. 25 . 59 . 24	
	22. 6		-	14	142	8.3				33. 9	22.3	*****	34	15	141	11	.0		20	

### CONCLUSIONS

The results of this investigation substantiate most of the conclusions arrived at in the 1932 investigation, modify two of the conclusions, and indicate some new conclusions. The conclusions substantiated are:

1. Materials of high specific gravity and their residues are, in general, more susceptible to changes in temperature than materials of low specific gravity and their residues.

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The following conclusions are somewhat modified from 1932:

Table 13.—Results of stability tests on cylinders exposed under plate and Vita glass

		Loss of	bitumen		8	Stability	at 77° F.	
Sample identi- fication	In 5	weeks	In 10	weeks	In 5 v	veeks	In 10	weeks
	Plate	Vita	Plate	Vita	Plate	Vita	Plate	Vita
	Percent	Percent	Percent	Percent	Pounds			Pound
1	. 2	1	2	1	225	250	275	37
2	8	5	8	7	850	700	850	80
13	7	4	6	5	450	450	500	42
14	4	2	4	3	475	400	500	45
17	13	12	15	13	650	700	650	70
18	18	16	18	17	500	500	525	- 50
19	8	6	8	8	1, 075	1,000	1, 250	1, 27
20	8	7	8	8	550	500	500	52
21	8	7	8	8	325	350	350	35
23	5	4	6	5	2:25	200	250	25
26	21	21	24	23	2,775	2,650	3, 550	3, 50
27	28	28	30	30	1, 225	1, 225	1, 575	1, 35
29	12	12	14	13	3, 775	3, 725	4,600	4.57
30	23	23	24	24	2, 575	2,550	2,850	3, 02
31	18	19	20	20	2, 325	2, 375	2,725	3, 12
32	18	17	19	20	1, 675	1, 725	2, 200	2, 67

5. The relative rates of volatilization of the various materials can be anticipated most readily from the distillation curves. The different classes of material may be differentiated in the loss and asphaltic-residue tests, especially if the time of reduction to 100 penetration is considered. However, sharp distinctions in initial curing properties, that may be of importance in some types of construction, can be determined only from the distillation curves.

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The following conclusions are developed upon the basis of the data collected in 1933 only:

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8. The use of Vita glass in place of plate glass for the cover of the exposure boxes did not materially change the results. However, because of the lateness in the year when these tests were started the results are considered inconclusive.

9. If like periods of the year are used for exposure, satisfactory check tests can be obtained with the exposure assembly used in these investigations.

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Table 14.—Comparison of 1932 and 1933 exposure tests

			Sam	ple ide	ntificat	tion		
Test characteristic	1	7	1	8	1	9	2	0
	1932	1933	1932	1933	1932	1933	1932	1933
Loss in 10 weeks, percent Penetration at 77° F Penetration at 32° F. Softening point, °F Ductility at 77° F., centimeters. Ductility at 34°–35° F., centi-	18. 3 41 17 163 1. 0	18. 7 36 14 170 0. 5	18. 5 132 37 106 110+	17. 7 152 43 105 110+	13. 3 40 9 121 110+	12. 0 44 10 121 110+	13. 1 95 30 115 65. 0	11. 5 136 41 112 90. 0
meters Insoluble in CS <sub>2</sub> , percent Insoluble in CCl <sub>4</sub> , percent Insoluble in 86° B. naphtha,	. 0 1. 06 8. 27	.0 3.17 9.66	1.5	7.5	.0 5.96 14.94	6.50 16.02	3.3	3. 8
percent	34. 8 19. 8	36. 2 19. 5 29	17. 6 18. 7 93	17. 3 19. 0 93	41. I 13. 7 28	38. 7 13. 8 29	26. 8 13. 7 67	24. 2 13. 6 84
Penetration at 32° F	12 181	12 168	29 113	28 116	8 129	128	24 121	28 120
meters Ductility at 34°-35° F., centi- meters	0.3	0.5	86.0	115.0	110+	110+	1.3	72.0
Insoluble in CS <sub>2</sub> , percent.	2, 50	3, 30			6.17	7.50		
Insoluble in CCl4, percent Insoluble in 86° B. naphtha,	9.11	9.88	. 17	.10	15. 61	18, 15	. 14	. 09
percent	37. 2	35.4	19.6	18.8	38.8	39.9	28. 5	27. 2
TESTS ON E	UBB.	ARD-	FIELI	CYL	INDE	RS		
Loss of bitumen:								
In 5 weeks, percent In 10 weeks, percent	15 18	15 19	18 19	16	10	7	9	12
In 15 weeks, percent Stability at 77° F.:1		21	21	20	14	11	13	13
Original cylinders, pounds. After 5 weeks' exposure,	275	150	200	125	325	175	225	100
pounds	1, 100	950	800	825	1, 375	1,650	775	725
pounds	1,575	1,000	1, 125	950	3, 175	2, 525	1, 525	1,000

<sup>1</sup> Differences in stability probably are caused by differences in grading of the sand.

1,550 1,200 1,650 1,200 4,050 3,075 1,975 1,250

## MOTOR-FUEL CONSUMPTION, 1934 [Compiled for calendar year from reports of State authorities]

												Classification	on of taxed	Classification of taxed motor fuel			
	Gross	Exemp	Exempted from pay- ment of tax 3	Shrink ance, d	Shrinkage allow- ance, discounts,	Gross	Subject to refund of entire tax	to refund	Amount		By rate of tax	of tax			By use a		Percentage change in
State	amount reported by State					assessed			on which tax was earned		At	At reduced rates	68 4	Doe blob	For ot	For other uses	from pre-
		Amount	Classes of use	Per- cent- age	Amount	ravarior	Amount	Classes of use		At full rate	Amount	Rate per gallon	Classes of use	way use 5	Amount	Classes of use	and control
	1,000 gallons	1,000 gallons			1,000 gallons	1,000 gallons	1,000 gallons	4	1,000 gallons	1,000 gallons	1,000 gallons	Cents		1,000 gallons	1,000 gallons		15.8
Alabama. Arizona. Arkansas. California	*75, 977 *75, 502 *134, 249 1, 356, 386	3, 944 3, 444 22, 361	E E		1, 276	70, 845 70, 845 129, 529 1, 320, 685	10, 280 3, 953 122, 030	NH, E	125, 576 125, 576 1, 198, 655	60, 565 108, 414 1, 198, 655	17, 162	6, 5, 4, 2	0	60, 565 119, 680 1, 198, 655	\$ 5,896	Z	13.0 4.3 2.1
Colorado Connecticut Delaware.	*172, 672 *308, 239 41, 556 246, 387	2,016 50,757 7,564	F, F.	62 1 183	3, 353	167, 303 254, 933 41, 556 235, 698	24, 013 6, 275 2, 042	NH, D, E	143, 290 248, 658 39, 514 235, 698	74, 381 248, 658 39, 514 235, 698	68, 909	*	ε	143, 290 248, 658 39, 514			2.5.00 4.5.00 4.5.00
Florida Georgia I daho I Illinois Indiana	243, 823 65, 828 1, 025, 751 465, 638	2, 203	24	8	2, 185	239, 435 63, 514 1, 025, 751 465, 638	5, 961 54, 877 26, 895	NNN	239, 435 57, 553 970, 874 438, 743	239, 435 57, 300 970, 874 438, 743	253	21/2	AV	57, 300 970, 874 438, 743	253	ΛV	25.5 4.6 8.7.
lowa. Kansas. Kentucky. Louisiana.	423, 886 *378, 781 184, 369 183, 977	7,366	F, NH, IC	60 64 60	12, 717 7, 575 5, 519	403, 803 283, 876 184, 369 178, 458	28, 805	H K	374, 998 283, 876 184, 369 178, 457	374, 998 283, 876 184, 369 178, 434	8	4	(1)	374, 998 283, 876			20.0 10.1 10.9 9.4
Marine Maryland Massachusetts Michigan	116,994 207,652 *590,625 735,593	1, 200	****			115, 794 206, 279 587, 828 735, 593	10, 616 21, 093 35, 763	F. NH NH NH	115, 794 195, 663 566, 735 699, 830	110, 924 193, 961 566, 735 698, 681	4,870 1,702 1,149	132	NH D	110, 924 13 195, 663 566, 735 698, 681	4,870	HN	20 20 E
Minnesota Mississippi. Missouri Montana	419, 454 130, 156 505, 677 *90, 567	1,908 2,180 2,562	F, E	00000	12, 534 2, 547 15, 170 2, 523	405, 012 125, 429 490, 507 85, 482	43, 500 11, 743 12, 211	HN HN HN	361, 512 125, 429 478, 764 73, 271	361, 512 112, 666 478, 764 73, 271	12, 763	1- 11	HZ	361, 512 112, 666 478, 784 73, 271	12, 763	HN	8. 16.5 5.4 33,2
Nebraska Nevada. New Jersey. New Jersey.	*224, 195 *27, 204 70, 652 *732, 761	2, 497	F, E, IC	60	6,726	216, 671 24, 707 70, 652 567, 839	2, 414 2, 352 2, 011	NN. NHHU	214, 275 22, 355 68, 641 567, 839	214, 257 22, 355 68, 641 567, 727	112	63	B	22, 355 68, 641 567, 727	112	B	28.5 28.5 16.8 3.9
New Mexico. New York. North Carolina. North Dakota.	*60,997 1,569,141 284,214 96,875	3, 922 52, 511 2, 540	F, E, IC F, P F, AV	21-1	898 15, 167 1, 877	56, 177 1, 501, 463 279, 797 96, 875	5, 043 37, 221 21, 485	HH NN	51, 134 1, 464, 242 279, 797 75, 390	51, 134 1, 464, 242 273, 686 75, 390	6, 111		Z	51, 134 1, 464, 242 273, 686 75, 390	6,111	X	12.9 11.2 17.6
Ohio 15. Oklahoma Oregon Pennsylvania 16.	*1, 303, 642 *312, 165 *165, 978 1, 136, 343	239, 900	F, E, IC	00 C1	31, 912 7, 985 22, 714	1, 031, 830 270, 432 165, 978 1, 113, 629	20,464	D, R NH, E	1, 031, 157 270, 432 145, 514 1, 113, 629	910, 214 270, 432 144, 917 1, 113, 629	120, 943		NH AV	910, 214 270, 432 144, 917	120, 943	NH AV	\$6.00 \$6.00
Rhode Island. South Carolina. South Dakota. Tennessee	*233, 167 130, 606 103, 122 207, 857	124, 303	F, E	-	4, 125	108, 864 130, 606 98, 997 201, 634	6,030	NH, E F D, R	102, 834 128, 646 98, 997 201, 627	102, 834 128, 646 89, 245 201, 627	9,752	52	BN	102, 834	9, 752	HN	9,3 5,6 8,7
Texas. Utah Vermont.	893, 802 64, 836 48, 550 264, 102	9,824	4	-8	8,840	875, 138 62, 858 48, 550 264, 102	84, 133	HN	791, 005 62, 858 48, 550 249, 540	791, 005 62, 858 48, 550 249, 540				791,005			11. 1 14. 9 10. 0 12. 6
Washington. West Virginia Wisconsin. Wyoming	260, 778 147, 610 431, 513 44, 111			23/2	10,788	260, 778 147, 610 420, 725 44, 111	21, 591 5, 217 35, 744	HEH	239, 187 142, 393 384, 981 44, 111	239, 187 142, 393 384, 981 44, 111				239, 187 142, 393 384, 981			10.1 15.8 1.5 25.5
District of Columbia.	147, 607	42, 186	F, F	C4	-	103, 820	169	NH		103, 129				103, 129			-0.9
Total	17, 220, 567	882, 618			201, 812	16, 136, 137	681,656		15, 454, 481	15, 210, 135	244, 346			12, 485, 421	162, 446		7.

B-Motor-boat use.
D-Fuel destroyed by fire, acts of God, etc.
R-Routine refunds (overpayment, etc.). The following symbols are used to designate certain classes or uses of motor fuel exempted from tax payment, subject to refund of the tax, or taxed at a lower rate:

NH—Uses other than for propelling motor vehicles on the highways. C—Use in public construction. F-Sales to Federal Government.
P-(Public) sales to State, county, or municipal governments.
F-Frue axported to other States or countries.
IC—Fruel moving through the State in interstate commerce.

 $A\,V-A\,v$  atton use. U-U se in vehicles licensed to operate exclusively in cities.

and for shrinkage, etc., and amount of motor fuel reported, prior to deduction of exempted fuel, allowance for shrinkage, etc., and amounts asblect to refund. Wheveve possible, fuel sempted because of export or
interstate movement, or to avoid duplication of tax payment, has been eliminated, in order that the total may
represent as closely as possible the total consumption of motor fuel in the State. Starred thems indicate that
unknown amounts of export sales or fuel moving in interstate commerce are included in the total shown

1 A number of States failed to report exempted fuel. Symbols are given only where amounts are reported.

2 Allowance for loss by leakage, evaporation, etc., and percentage its computed on the gross
reported; in others on the net after deduction of exempted fuel. In some States the percentage is fixed; in others
it is the maximum allowable.

The purpose of this castification is of situation and remained to taxed motor fuel by motor
whicles operating on the highways, and consumption for other purposes. In the case of states which do not
make this distinction, the classification is omitted.

The purpose of this castification is omitted.

The purpose of the sale classification is omitted.

The purpose of the classification by use. In these cases the percentage is absent on the total amount away on the total amount on which tax was
earned. The total on which the Nation-wide percentage is based on the total amount on which tax was
earned. The total on which the Nation-wide percentage is based on the decrease (District
of Columbia only) is indicated by a minus sign.

7 Refunds on nonhighway use not allowed after Feb. 12, 1934.
7 A12,000 gallons at 6earls prior to Feb. 13, 1934.
7 A12,000 gallons at 6earls prior to Feb. 13, 1934.
Factors at 4 earls, 8,577,000 gallons; at 2 cents, 935,000 gallons.
1 Estimated by State
10 Actual allowators perforted: no faced percentage.
10 Actual allowators reported: no faced percentage.
11 Rate of 4 cents per gallon applies to any gas-generating liquid having a flash point below 110° F. Additional.
11 Rate of 4 cents per gallon applies to any gas-generating liquid having a flash point below 110° F. Additional.
12 Refunds are made on all nonhighway uses with the exception of fuel used in commercial motor boats.
13 A Scent tax is imposed on motor-vehicle their and a 1-cent tax on all liquid fuels commercial and kerosene.
14 A Scent tax is maposed on motor-vehicle their and a 1-cent tax on all liquid fuel reported was 1,205,642,000 gallons.
16 Tax is imposed on all liquid fuels, including fuel oil and kerosene, usable in internal-combustion engines.

### STATE MOTOR-FUEL TAX EARNINGS, 1934

[Compiled for calendar year from reports of State authorities 1]

	Tax						Cla	ssificatio	n of tax e	arnings 1		Oth		ings in o		on	
	per g	allon		Gross	Re-	Net earn- ings	Ву	rate of t	ax	By use o	f fuel 2		with m	iotor-fue	l tax 3		Grand
State	On	On	Date of rate change	tax as- sessed 4	funds earned or paid *	on all motor- fuel		At reduc	ed rates <sup>6</sup>	For	For	Dis- tribu-	Deal-	In-	Other		total earn- ings
	Jan.	Dec.			,	taxed 4	At full rate	Rate per gallon	Amount	high- way use <sup>2</sup>	other uses	tors' li- censes	ers' li- censes	spec- tion fees	fees, etc.	Total	
labama	Cents	Cents 6		1,000 dollars 9, 299	1,000 dollars	1,000 dollars 9, 299	1,000 dollars 9, 299	Cents	1,000 dellars	1,000 dollars	1.000 dollars	1,000 dellars	1,000 dellars	1,000 dollars	1,000 dollars	1,000 dollars	1,000 dollars 9, 2
rizonarkansas alifornia	5 6 3	5 63/2 3	Feb. 3	3, 542 8, 118 39, 621	514 254 3, 661	3, 028 7, 864 35, 960	3, 028 7, 047 35, 960	6, 5, 4, 2	7 817	3, 028 5 7, 481 35, 960	* 383	10			1	10	3, 0 7, 8 35, 9
olorado	9.4	4	Feb. 1 and Sept. 1.	7, 591	1, 116	6, 475	3, 719	8.4	2,756	6, 475							0.4
Connecticut Delaware Plorida	2 3 7	3 7		5, 099 1, 246 16, 499	126 61	4, 973 1, 185 16, 499	4, 973 1, 185 16, 499			4,973 1,185			11 35	******		35	4, 90 1, 18 16, 50
Georgiadahollinoisndiana	6 5 3 4	6 5 3 4		14, 366 3, 169 30, 772 18, 626	298 1, 646 1, 076	14, 366 2, 871 29, 126 17, 550	14, 366 2, 865 29, 128 17, 550	23/2	6	2, 865 29, 126 17, 550	6				1 5 20	1 5 20	14.36 2.87 29.17 17.57
owa Kansas Kentucky Jouisiana	3 3 5 5	3 3 5 5		12, 114 8, 516 9, 218 8, 923	864	11, 250 8, 516 9, 218 8, 923	11, 250 8, 516 9, 218 8, 922	4	i	11, 250 8, 516		5		76	37 3	118 3	11. 2 8. 6 9. 2 8. 9
Maine Maryland Massachusetts Michigan	4 4 3 3	3		4, 632 8, 251 17, 635 22, 068	146 442 633 1,090	4, 486 7, 809 17, 002 20, 978	4, 437 7, 758 17, 002 20, 961	1 3	49 51 17	12 7, 809 17, 002	49	******	4		2	6	4. 4 7. 8 17. 0 20. 9
Minnesota Mississippi Missouri Montana	3 6 2 5			12, 150 7, 519 9, 810 4, 274	1, 305 631 235 610		10, 845 6, 760 9, 575 3, 664	1	128	. 10, 845 6, 760 9, 575 3, 664		*****	1	97	13 137 8	98 137 106	10,5 7,6 9,6 3,6
Nebraska Nevada New Hampshire New Jersey	4 4 4 3	4		8, 667 988 2, 826 17, 035	97 94 86	894 2,746	8, 570 894 2, 746 17, 032	2		894 2,746 17,032			25	85	1	86	8. 2. 17.
New Mexico New York North Carolina North Dakota		3 6		2, 809 45, 044 16, 788 2, 906	252 1, 117 306 644	43, 927 16, 482	2, 557 43, 927 16, 421 2, 262	1	6	2, 557 43, 927 1 16, 421 2, 262	61	. 9	58			18 58 709	2,1 43,1 17, 2,1
Ohio Oklahoma Oregon	-	4 4		38, 982 10, 817 8, 299		10, 817	36, 409 10, 817 7, 246		1, 20	10, 817		-				4	37, 10,
Pennsylvania		3 3		. 33, 409		33, 409	33, 409	*****		6 7, 246						4	33,
Rhode Island South Carolina South Dakota Tennessee		2 2 6 6 4 4 7 7		2, 177 7, 836 3, 960 14, 114	11 19	7 7,719	2, 057 7, 719 3, 570 14, 114	2	19	5 3, 570				3			7. 3. 14.
Texas Utah Vermont Virginia		4 4 4 4 4 5 3		35, 003 2, 514 1, 943 13, 203	3, 36	5 31,640 2,514 1,942	31, 640 2, 514 1, 943			31, 64				1	11	1 111	31, 2, 1, 12,
Washington West Virginia Wisconsin Wyoming		5 4 4 4 4 4 4 4		13, 031 5, 90 16, 82 1, 76	9 1,08 5 20 9 1,43	11, 959 9 5, 696 10 15, 399 1, 764	11, 956 5, 696 15, 399 1, 764	3		11, 95 5, 69 15, 39	9			5		1 8	11, 5, 15,
Dist. of Columbia.  Detailed totals 10	-		2	2, 07	7 1	2, 063				2, 06	3						2,
Grand totals			17 3, 66	591, 99	5 26, 96	38 565, 027	559, 72	9	5, 21	434, 63	2, 05		0 14	1 1,09		5 1.615	566,

¹ See preceding table for gross gallons of motor fuel reported, exemptions, allowances, etc., gross gallons taxed, gallons subject to refund, net gallons taxed, and information regarding classes of use exempted, subject to refund, or taxed at lower rates.

² The purpose of this classification is to distinguish between the tax earnings on motor fuel sold for use in motor vehicles on the highways and tax earnings on motor fuel sold for other uses. In the case of those States that do not make this distinction, the classification is omitted.

³ Amounts less than \$500 not tabulated.

¹ In the great majority of cases the assessments or earnings of the calendar year were reported. A few States reported the actual collections of the year, which lag the assessments by 1 to 2 months.

¹ In most cases the refunds reported were those actually paid during the year, rather than refunds claimed on motor fuel purchased during the year. The error involved in deducting refunds paid from gross tax assessed tends to balance over an annual period. The refunds tabulated include both refunds of the entire tax and partial refunds.

¹ In the case of Arkansas and Colorado, where the rate was changed during the year, the tax earnings at the lower rates, 6 and 4 cents, respectively, are shown under this heading.

<sup>7</sup> Includes \$445,000 on 6-cent tax prior to Feb. 13, 1934, and amounts at reduced rates at State borders, as follows: At 5 cents, \$7,000; at 4 cents, \$347,000; at 2 cents, \$18,000.

<sup>8</sup> Estimated by State.

<sup>9</sup> Rate was 5 cents from Feb. 1 to Aug. 31, 1934.

<sup>10</sup> Retail gasoline station licenses, \$45,000, included in report on motor-vehicle receipts.

<sup>11</sup> Includes distributors' licenses.

<sup>12</sup> Refunds are made on all nonhighway uses with the exception of fuel used in commercial motor boats. Earnings on motor-boat fuel (amount not reported) are included.

<sup>13</sup> Includes \$138,560, earnings on special gasoline tax collected in Gulf Coast counties (Hancock, Harrison, and Jackson) for seawall protection, and \$1,559 in penalties, less \$2,629, refunds for notary fees.

<sup>14</sup> Inspection fees on gasoline and kerosene; bulk of receipts on paroline.

<sup>15</sup> Includes dealers' licenses.

<sup>16</sup> Classification by use includes 36 States and the District of Columbia.

## AS PROVIDED BY SECTION 284 OF THE NATIONAL INDUSTRIAL RECOVERY ACT (1934 FUNDS) AND BY THE ACT OF JUNE 18, 1934 (1935 FUNDS) CURRENT STATUS OF UNITED STATES PUBLIC WORKS ROAD CONSTRUCTION

CLASS 1.—PROJECTS ON THE FEDERAL-AID HIGHWAY SYSTEM OUTSIDE OF MUNICIPALITIES

AS OF JULY 31, 1935

4 40	APPOL	APPORTIONMENTS		COM	COMPLETED			UNDER CONSTRUCTION	TRUCTION		APPROVED	APPROVED FOR CONSTRUCTION	JCTION	BALANCE OF FU	BALANCE OF FUNDS AVAILABLE
SIATE	Sec. 204 of the Act of June 16, 1933 (1934 Fund)	33 June 18, 1934 (1935 Fund)	34 Total Cost	1934 Public Works Funds	1935 Public Works Funds	Mileage	Estimated Total Cost	Public Works Funds	Public Works Funds	Mileage	1934 Public Works Funds	Public Works	Mileage	FOR NEW 1934 Public Works	V PROJECTS 1935 Public Works
Alabama Arizona Arkansas	3,947,753	2,129,921 1,338,712 1,714,000	\$6,527,896 112 5,076,955 00 3,828,359	356 3,460,745 3,829,744 359 2,576,551	139,902	358.2	2,147,304	\$ \$60,140	1,364,071	32.2	20,073	195,768	11.8	6,795	# #30,180
California Colorado Connecticut	7,912,928	3,713,643 2,424,504 3 607,500	11,199,998 04 5,475,319 00 1,135,531	7,789,341 3,357,026 31,109,185	1, 609, 765 1, 895, 044		3.163,930 625,115	122,227	2,122,000		505.65	906,903	22.8	1,360	74,974
Delaware Florida Georgia	2,469,370 5,045,592	1,116,600 2,556,745		868.470 15 2.355.070 69 3.977.140	271,538		149.545	66,825	140,992		578	505.345	50 t	23,507	4.589 33.068
Idaho Illinois Indiana	2,166,858 4,408,827 5,018,921			14 2,027,114 89 2,391,815 62 4,243,283	363,290		681,542 4,088,825 3,018,635	2,002,682	2,086,143	39.7	3,000	1,670	100 m	7,986	741,845
Iowa Kansas Kentucky	5,027,830 5,044,802 3,751,605	2,354,131	5,506,127 5,933,599 3,856,210	27 4,978,830 99 5,004,394 10 3,408,445	796,564	331.3	2,002,083 1,630,648 1,398,252	31,530	1,555,141	125.1		13,700	a) n	275.83	254,000
Louisiana Maine Maryland	2,693,135		9 2,864,278 5 1,890,999 9 916,280	78 2,399,450 39 1,541,605 10 791,495	93,648 319,255 123,550	78.3	1,434,550 472,624 1,030,226	234,221 11,351 900,555	1,047,066	27.3	51,291		2 2	8,173	239,706
Massachusetts Michigan Minnesota	1,101,716 6,051,533 4,561,011	1,582,874 3,226,284 2,533,733	5,670,800	7 1,048,966 0 5,213,496 2 4,266,504	1,740,036	242.3	3,441,400	52,687 782,950 217,577	2,637,386	145.9	60.000	313,975	410	55.067	587.503
Mississippi Missouri Montana	3,489,337	2,832,162	5,647,003	3 2,682,221 0 4,409,896 2 4,345,161	232,942	292.9	2,563,432	723,191	1,276,205	78.2	23,588	776.607	28.00	60,337	25,971
Nebraska Nevada New Hampshire	3,914,461	1,982,182	5,154,215 3,442,296 937,865	5 3,859,875 2,666,269 5 692,118	216,519 703,321 201,292	378.1	2,054,603 884,019 243,690	21,644 204,818	1,688,363	147.2	17.734	75,122		2,963	7,528
New Jersey New Mexico New York	3,173,019 2,846,648 10,465,672	1,676,769	2,247,203	2,025,846	1,047,273	374.7	1.770.707 631.786	1,113,074	\$60,860 629,496 3,027,452	11 S. C. L.	11,623	10.167		34,100	350,605
North Carolina North Dakota Ohio	4,761,147 2,902,224 7,277,758	1,469,464	5,168,217	3,684,500		1,165.4	1,435,381	663,506	565,944 324,435	178.6	91,438	322,565	21.1	321,703	169,549 518,146 677,556
Oklahoma Oregon Pennsylvania	4,608,399 3,053,448 6,691,194	2,342,590	5,005,045 3,584,581 7,956,871		.,	330.6	2,000,652		1,436,273	70.7	6,016	143,167	7.0	13,669 4,272 45,196	593,305 146,499 69,449
Khode Island South Carolina South Dakota	2,729,563	474,772 940,954 1,523,821		899,627 2,359,621 2,532,214	168,738 214,683 493,190	223.4	391,854 51,548 1,285,130	79.740	295,834 275,918 809,360	2,000 2,000	18,520	51,576	0,0	35,811	10,200
Texas Utah	4,246,309 11,588,643 2,367,205	2,105,454 6,858,253 1,066,345	5,226,315	3,999,811	1,503,032	1,161.1	1,508,309	176,547 247,933 37,000	1,235,361 4,226,173 341,550	338.23	35,675	183,445	0.27	67.75	109,304
Vermont Virginia Washington	928,184 3,731,207 3,057,934	1,916,178	1,135,574 4,340,679 3,329,879	912,376	124,795 632,668 474,277	183.1	356.137 1,447.575 1,266,948		1,043,045	14.1	4,042	3,866	1. 9.	11,766	6.333
Wyoming District of Columbia	2,013,405	1,140,167	2,301,844	1,900,800	359,092 361,138 619,195	238.6	496,108 1,707,503 1,095,104	82,643 273,603 141,021	413,465	14.8	27,869	47,222 69,796 43,808	3.5	2,093 6,403 4,503	320,389
Hawaii	1,693,344	598,778	\$29,029	526.724		19.3	1,430,996	1,126,854	454.66	20.8	20,973	183,392	2.1	18.791	2115 363
TOTALS	185,235,236	93.935,660	213.731,417	165,756,136	20 744 ASO 4C					-				171111	300000

# CURRENT STATUS OF UNITED STATES PUBLIC WORKS ROAD CONSTRUCTION

AS PROVIDED BY SECTION 204 OF THE NATIONAL INDUSTRIAL RECOVERY ACT (1934 FUNDS) AND BY THE ACT OF JUNE 18, 1934 (1935 FUNDS)

CLASS 2.—PROJECTS ON EXTENSIONS OF THE FEDERAL-AID HIGHWAY SYSTEM INTO AND THROUGH MUNICIPALITIES

AS OF JULY 31, 1935

	APPORTI	APPORTIONMENTS		COMPLETED	TED			UNDER CONSTRUCTION	TRUCTION		APPROVED	APPROVED FOR CONSTRUCTION	CTION	BALANCE OF FU FOR NEW	BALANCE OF FUNDS AVAILABLE FOR NEW PROJECTS
BTATE	Sec. 204 of the Act of June 16, 1933 (1934 Fund)	Act of June 16, 1934 (1935 Fund)	Total Cost	1934 Public Works Funds	Public Works Funds	Mileage	Estimated Total	Public Works Funds	Public Works Funds	Mileage	Public Works Funds	Public Works Funds	Mileage	Public Works Funds	1935 Public Works Funds
Alabama Arizona Arkansas	8,389,928 756,982 1,964,534	\$ 1,064,961 305,191 657,025	\$2,000,340 738,683 1,902,627	\$1.872.02\ 622.80\ 1.591.572	\$ 119,375 86,475 210,081	15.1 15.1 15.3	# 786.135 323.930 510.198	\$ 463,060 129,322 246,967	\$ 323.075 156.833 259.979	33. 15. 8 14. 15. 15.	\$ 29.878	\$ 122,470 311,035	3. 2.	\$ 24.965 1,857 22,913	\$ 500.041 61,883 75,931
California Colorado Connecticut	4,213,986 1,718,633 802,407	2,219,360 190,000 426,500	5,358,835 1,937,435 838,549	3.849.724 1.672.058 802,407	847,032 169,411 9,362	65.6 40.4 10.2	2,680,138 11,229 193,692	356.840	1,070,354	8.3		194,200	5.5	7,422	107,784 20,589
Delaware Florida Georgia	1,459,648	230,849 501,200 1,276,373	542,664 1,839,561 2,414,917	1,459,771 2,232,680	73.875 102,827 146,915	21.4	18,165	275.754	163,804	1.1		139,600	1.2	17,554	138,809 94,969 693,715
Idaho Illinois Indiana	1,197,629 7,381,910 4,287,050	321,126 2,230,350 2,246,858	1,222,156 6,593,153 3,615,513	1,151,479 6,285,871 3,362,976	27,061 8,850 104,496	8.35 6.24	190,673 2,121,438 2,280,824	1,036,345	189,485	 39.0	147,260	35,000 620,357 329,769	82.5	12,435 27,549	69.560 516.050 282,342
Iowa Kansas Kentucky	2,614,472 2,522,401 1,927,628	1,311,000 1,432,949 956,599	2,303,093	2,105,968 2,473,462 1,476,834	101,635 377,503 71,226	3.55	1,406,378	908.W65	808,800 1,044,466 528,141	23.1 11.3 9.9	30,000	10,980	- 35	31,436	262,865
Louisiana Maine Maryland	1,706,577	744,560 484,379 452,585	946,632 925,109 390,021	793,980 872,233 384,017	147,468	12 to 10 to	1,197,913	896.730 83,155 262,457	259,141	300	10,666	203,661	000	7,200	134,271 6,907 452,515
Massachasetts Michigan Minnesota	5,007,199 3,500,637 3,719,143	847,600 1,613,142 1,421,494	3,392,363	2,102,182 3,118,335 3,136,040	96.359 133.100 376.049	113.9	3.071.272	342,450	1,262,400	17.2	19,400	130.975	3.03	28,646	579,317 86,667 415,538
Mississippi Missouri Montana	1,744,669	354,082	1,166,465	1,033,466 3,001,268 1,016,328	214,956 24,854 49,236	88.3	1,142,578	626,097 867,762 64,410	120,417	123.0	36.943	18,467	0 M4 0.40	150.471	100,163
Nebraska Nevada New Hampshire	1,957,240 500,051 740,335	991,091 100,000 242,465	5,498,414 5,39,562 845,980	1,915,583	546.276	10.3	53,951		53.741	2.4	26,150 13,044	141,402	9.7	41.657	146.743
New Jersey New Mexico New York	3.117.921 1.674.158 8.255.661	1,809,500	3,060,011	2,810.715	106,606 150,097 364,600	25.9 40.2 63.4	1,225,905	182,690	804,667 69,432 2,964,090	5.9	1,010	631,441 183,963 219,150	9-4	17,794	264.787 96.014 393.850
North Carolina North Dakota Ohio	2,380,573	1,210,236	2,791,849	2.135,990 1,241,533 4,238,314	634,661 92.515	96.00 9.00 9.00 9.00	946,276	103.576 99.844 93.352	426.700 88.185 1,374.172	4.000	91,217	91,264	16.4	49,790 22,366 4,020	363,480
Oklahoma Oregon Pennsylvania	2.304.200 1.526.724 4.854.988	2,397.703	2,415,796	2,112,039 1,415,650 4,279,869	220,827 150,853 725,785	72.7	786.161 619.349 1.354.210	192,161 66.635	575.152	0.63	518	126,468 80,000 285,851	6.00	43,921 63,976	248,847 92,418 633,092
Rhode Island South Carolina South Dakota	1,564,791	285,760 4468,000 761,911	1,236,620	518,991 1,199,812 1,188,674	36.001 23.620 43.905	38.0	105.759	156.274	106.759 164.747	9,50	8.705 60.45	15,412	4.7	60.634	197,422
Tennessee Texas Utah	2,123,155 6,642,863 776,826	1,121,789	5,209,866 6,049,343 612,144	1,921,523	264,516 216,679 97,066	124.6 20.8 80.8	6%6,283 1,662,930 650,283	201.632 671.637 129.130	494.651 814,902 433,800	555	16,465	115,946 398,296	8.7	246,459	246.675 365.123 2,306
Vermont Virginia Washington	500,509 1,948,780 1,977,260	240,611 956,021 776,603	636,840 2,866,435 2,330,294	1,922,046	96,285 352,905 365,313	2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	137,402 702,604 412,559	26,801 246,442 53,369	105,062 396,439 359,190	5.03	13.975	39.244 155.506	6.5	15.036 7.002 1.845	51.170
West Virginia Wisconsin Wyoming	1,342,270	1,379,513	3,108,178	2,509.104	28,109 514,579 2,784	62.2	494,743 629,047 162,381	270.594	629.047	10 00 m	28.642	47.791 215.499	900	29,375	20.035
District of Columbia Hawaii	946,4415	121,051	877,332	196,781	121,051	6.9	250,16%	250,164		Pa					200
TOTALS	115,617,401	M8.020,938	111,510,588	97,431,410	9,404,416	2,021.2	41,300,530	15,058,236	22,570,806	453.2	864,143	6,806,844	104.7	20.36.8 £255	N alla Alla

## AS PROVIDED BY SECTION 204 OF THE NATIONAL INDUSTRIAL RECOVERY ACT (1934 FUNDS) AND BY THE ACT OF JUNE 18, 1934 (1935 FUNDS) CURRENT STATUS OF UNITED STATES PUBLIC WORKS ROAD CONSTRUCTION

CLASS 3.—PROJECTS ON SECONDARY OR FEEDER ROADS

STATE	APPORT	APPORTIONMENTS		00	COMPLETED		-		UNDER CONSTRUCTION	Tana and a second						
21610	Sec. 204 of the Act of June 16, 1933	Act of June 18, 1934	Total Cos	-	-			-		MOCITON		APPRO	APPROVED FOR CONSTRUCTION	TRUCTION	BALANCE OF FOR NE	BALANCE OF FUNDS AVAILABLE FOR NEW PROJECTS
4	(May Fund)			Funds Funds	Public Works Funds		Mileage	Estimated Total	Public Works	Public Works Funds	Mileage	Public Works Funds	1935 Public Works Funds	ks Mileage	1934 Public Works	1935 Public Works
Arizona Arkansas	2,032,452 599,427 1,449,634	1,064,960 998,032 857,024	2,016,508 2,985,299 1,382,268	530,962 1,249,425	•	357.034	24.3.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.	622.670 648.037	187,785	634,885	9.3		\$ 128,940	10.6	* 141. sta	-
Colorado Connecticut	3,480,440	1.999.203	2,601,784	-		-		833.84t	323.897	1,364,725			229, 946		53,461	77.009
Delaware Florida	481,113	230.849	-			-	-	222,880	110,000	320,140	39.5		206.527	4.6	1,700	155,951
Georgia	2,320,973	1,043,543	1,626,676	1.590,474	331,460		57.1 68.5	265,716 704,065 060,465	215,448	41,690	2 19		216 212			185,099
Illinois Indiana	5.740,033	4,262,273 151,472	3.633, 423	3,495,530	396.344	44 (0	10		265,472	247,061				-	103,711	785,690
Iowa Kansas Kentucky	2,522,401	1,330,595				-			274,972	103,626	50.2		473.156		13,058	334,400
Louisiana Maine	1,426,879	838,953 Mile 913						1,341,513	305.453	1,211,984	167.8	43,499	29,555	2,55	707	9
aryiand	425,185	1.067.934		865,179	390,302		66.5	59.505	16,123	589,699	38.7	130,522	129.772		374	18,305
Minnesota Mississippi	3,184,057	1,613,142	3.275.590	2.937.940 2.175.788	103,450	207.7		1,672,663	205,227	1,467,417	5.5	200	313.670	8. 0°.	30,44,81	425,574
Missouri	2,923,273	2,363,922	3.194.579	2,636,006	16.500	143.1				85,852	1.60 of		112,651	-	51,133	15.866
Nebraska Nevada New Hampshire	1,156,479	991,091	2,472,284	1,957,240			-		-	311, 474	31.9	82,153	55,435	29.0	1,716	3.1%
New Jersey New Mexico New York	55,099	460.000	570,258	55,099					79,000	180,528	6.4	8,443	169,589	15.2	2,683	19.581 54.195 12.181
North Carolina North Dakota	2,380,573	3,693,000	2,865,726	3.027.536					36.931	351,820	41.14	65,732	170.412	1.1		182,063
Ohio	3.871.148	1,966,253	4,172,803	3.751.018	169,850	387.3	_	1.023,042	150.265	50,717	54.6	100 301	134,315	9.6	43.584	15, 104
Ostanoma Oregon Pennsylvania	2,304,199 1,526,724 7,344,822	1.171.295	2,304,496	2.052,553	70.199		-		-	1,318,823	89.8		27,200	0.00	28,938	328,960
Rhode Island South Carolina South Dakota	1,364,716	1,342,000	1,214,563	439.716 1.086.655	341,038	33.1		get.			198.5		137.603	7.0	12,317	174,883
Tennessee	2,123,155	1,075,748		1,325,467	131,244	136.9	_				167.5		36,815	1.4	14,619	4,663
Utah	1,048,677			5,952,792	694,201	872.7	2,911,390		34,517 2,	2,866,130 295,000	37.6	17,894	121,139	5.1	63,978	221,404
Virginia Washington	1,736,770	893,188	1,716,601	1,551,099 1,080,673	166,042 105,009 267,589	216.3	80,229		112,886		5.3	sh Gun			3,019	1,000
West Virginia Wisconsin Wyoming	1,118,559 2,431,220 1,125,332	1,743,354	2,644,458	2,151,288	308,635	179.7	-				22.7		470,075	15.0	16,141	1,264
District of Columbia				971,729	135,376	156.8	450,3		75,284		56.7		123,338	220	2.590	115,768
TOTALS	-	-	-	177,718 81.877 Mas		4.9		50		393,065	2.5		75.395	9.4	562	135.097
		_	_	2011101	11,996,937	10,021,3	48.601 412	22 0 00 0	-		-		11.	200		169,347

### PUBLICATIONS of the BUREAU OF PUBLIC ROADS

Any of the following publications may be purchased from the Superintendent of Documents, Government Printing Office, Washington, D. C. As his office is not connected with the Department and as the Department does not sell publications, please send no remittance to the United States Department of Agriculture.

### ANNUAL REPORTS

Report of the Chief of the Bureau of Public Roads, 1924. 5 cents.

Report of the Chief of the Bureau of Public Roads, 1927. 5 cents.

Report of the Chief of the Bureau of Public Roads, 1928. 5 cents.

Report of the Chief of the Bureau of Public Roads, 1929. 10 cents.

Report of the Chief of the Bureau of Public Roads, 1931. 10 cents.

Report of the Chief of the Bureau of Public Roads, 1932. 10 cents.

Report of the Chief of the Bureau of Public Roads, 1933.

Report of the Chief of the Bureau of Public Roads, 1934.

### DEPARTMENT BULLETINS

No. 136D . . Highway Bonds. 20 cents.

No. 347D . . Methods for the Determination of the Physical Properties of Road-Building Rock. 10 cents.

No. 583D , . Reports on Experimental Convict Road Camp, Fulton County, Ga. 25 cents.

No. 1279D . . Rural Highway Mileage, Income, and Expenditures, 1921 and 1922.

### TECHNICAL BULLETINS

No. 55T . . . Highway Bridge Surveys. 20 cents.

No. 265T . . . Electrical Equipment on Movable Bridges. 35 cents.

### MISCELLANEOUS CIRCULARS

No. 62MC . . Standards Governing Plans, Specifications, Contract Forms, and Estimates for Federal-Aid Highway Projects. 5 cents.

### MISCELLANEOUS PUBLICATIONS

No. 76MP . . The results of Physical Tests of Road-Building Rock. 25 cents.

Federal Legislation and Regulations Relating to Highway Construction. 10 cents.

Supplement No. 1 to Federal Legislation and Regulations Relating to Highway Construction.

No. 191 . . . . Roadside Improvement. 10 cents.

The Taxation of Motor Vehicles in 1932. 35 cents.

### REPRINT FROM PUBLIC ROADS

Reports on Subgrade Soil Studies. 40 cents.

Single copies of the following publications may be obtained from the Bureau of Public Roads upon request. They cannot be purchased from the Superintendent of Documents.

### SEPARATE REPRINT FROM THE YEARBOOK

No. 1036Y . . Road Work on Farm Outlets Needs Skill and Right Equipment,

### TRANSPORTATION SURVEY REPORTS

Report of a Survey of Transportation on the State Highway System of Ohio (1927).

Report of a Survey of Transportation on the State Highways of Vermont (1927).

Report of a Survey of Transportation on the State Highways of New Hampshire (1927).

Report of a Plan of Highway Improvement in the Regional Area of Cleveland, Ohio (1928).

Report of a Survey of Transportation on the State Highways of Pennsylvania (1928).

Report of a Survey of Traffic on the Federal-Aid Highway Systems of Eleven Western States (1930).

A complete list of the publications of the Bureau of Public Roads, classified according to subject and including the more important articles in PUBLIC ROADS, may be obtained upon request addressed to the U. S. Bureau of Public Roads, Willard Building, Washington, D. C.

## CURRENT STATUS OF UNITED STATES PUBLIC WORKS ROAD CONSTRUCTION

AS PROVIDED BY SECTION 204 OF THE NATIONAL INDUSTRIAL RECOVERY ACT (1934 FUNDS) AND BY THE ACT OF JUNE 18, 1934 (1935 FUNDS)

### SUMMARY OF CLASSES 1, 2, AND 3.

AS OF JULY 31, 1935

	APPORTIC	APPORTIONMENTS		COMPLETED	TED			UNDER CONSTRUCTION	TRUCTION		APPROVED	APPROVED FOR CONSTRUCTION	CTION	BALANCE OF FU	BALANCE OF FUNDS AVAILABLE FOR NEW PROJECTS
STATE	Sec. 204 of the Act of June 16, 1933 (1934 Fund)	Act of June 18, 1934 (1935 Fund)	Total Cont	Public Works Funds	Public Works Funds	Milesge	Estimated Total	Public Works Funds	1935 Public Works Funds	Mileage	Public Works Funds	1935 Public Works Funds	Mileage	1934 Public Works Funds	Public Works Funds
Alabama Arizona Arkansas	\$ 8,370,133 5,211,960 6,746,335	2,541,935	8 10,544,744 6,801,137 7,113,454	7,135,888	\$55,986 1,006,887 965,678	550.3 1666.5	\$ 3.756.109 1.874.076 2.823,623	\$ 1,110,985 174,322 1,096,989	2, 322, 031 1, 996, 161 1, 503, 655	165.9 78.5 153.6	\$ 49,951 15,039 140,639	\$ 447,176 770,563	39.1	\$ 73,309 39,090 93,159	\$ 1,034,647 138,892 188,133
California Colorado Connecticut	15,607.354 6,874.530 2,865.740	7.932,206 3,486,006 1,454,868	20,609,040 10,014,539 2,671,380	14,793,906 6,637,716 2,570,712	1,672,796 2,615,817 22,25	575.4	7,677,909 1,167,154 1,553,894	802.964 198,405 295,028	4,557,069 836.831 920,127	70.5		1,363,631	37.7	10,1462	338.709
Delaware Florida Georgia	1,819,068 5,231,834 10,091,185	923.395 2.661.343 5.113.491	2.207.097 6.903.853 9.067.538	1.594.418 5.074.597 7.800.494	576.536 705.845 971.664	235.6 539.0	433,426 1,581,213 3,628,865	223,925 69,148 1,819,406	1,426,517	18.5 72.7 176.9	23,966 6,394	378,683	7.8	167 64,121 464,591	146,013
Idaho Illinois Indiana	17.570.770 17.570.770 10.037.843	2.277,486 8.921,401 5.088.963	5.355.529 12,809,564 8.500,128	4.273.123 12.173.189 8.027.933	786.715 245,922 157.034	423.4 311.8 243.3	1.119.277	131.759 5.304.498 1.658.495	954.290 6.554.131 3.965.895	67.2 364.9 223.8	50,260	36,670	11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	81,367 42,823 125,350	497,811 850,450 397,732
Iowa Kansas Kentucky	10,055,660	5,118,361 5,117,675 3,818,311	10,610,864 11,266,962 7,639,339	9.378.197 9.551.326 6.657.329	928,005 1,323,424 543,581	915.3	4,966,043 4,177,107 3,685,737	336.382	3.505.891	214.1	92,417	167,600 37,222 190,281	**** ****	8.878 806.735	516.865
Louisiana Maine Maryland	5.828.591	2.963.932 1.711,586 1.810.058	5,011.374 4,162,421 2,280,231	3,256,241	342,291 756,964 163,680	3 E.S.	3,432,133 892,890 2,461,446	1,340,921	1,895,906 784,708 468,790	18.2	192,479	333.452 158.253 222.970	100 mg	15.747	392,282 11,661 914,478
Massachusetts Michigan Minnesota	6,597,100	3,350,474 6,452,568 5,425,551	4,200,967	3,620,889	98.359	67.6	4,694,815 6,756,794 2,957,995	2,929,059	1,686.958 5,367.203 1,903,508	141.0 247.9 216.3	19,400	313,670	15.4°5 10.7°5	47,152 116,430 133,835	1.249,487 217.939 457.375
Mississippi Missouri Montana	6,978,675 12,180,306 7,439,748	3,540,227 6,173,740 3,769,734	8,078,785 11,550,962 9,998,341	4,964,483 10,097,170 7,102,183	820,235 689,581 2,389,133	473.0 943.8 897.3	5,682,707	1,809.872 1,604,996 68,540	1,462,473	206.1 450.1 96.6	60.531 245,950 147,641	1,666,261	89.3 1.69.3	143,786	98,788
Nebraska Nevada New Hampshire	7,828,961 4,545,917 1,909,839	3,964,364	10.12%,912 5.569,952 2.35%,103	1,762,697	1,263,400	53.6	2.756.165	21.644 216,818 29,000	2,389,925 806,830 468,508	175.7	52.327	112,536 224,197	17.0	23,249 23,249 56,515	198.502 62,479 51.056
New Jersey New Mexico New York	6,346,039 5,792,935 22,330,101	3,220,879 2,941,700	5,383,742	4,891,660 5,665,578 19,340,340	136,352	672.9	3,104,136	36,931	1,473,052	95.9	12,633	812,020 183,963 238,050	1.1	158,615	797.455 108.232 644.061
North Carolina North Dakota Ohio	9,522,293 5,604,446 15,464,592	2,936,967 7,865,012	10, 828, 792 5,986, 728 17,011, 922	7.997.214 5.055.276 15,026,171	1,746,315	1.061.5	3.006,699 926,427 6,284,657	927.348 325.115 370,463	1,955,421	242.24 243.2 165.4	331.961	548,144 651,058 107,800	32.8 191.7 6.5	415.077 92.493 87.958	591,061 1,369,996 1,514,871
Oklahoma Oregon Pennsylvania	9.216.798 6.106.896 18.891,004	4,685,180 3,097,814 9,590,788	9,725,336 7,316,249 19,863,931	8,414,407 5,816,229 16,945,385	907.677 877.871 2.153.200	653.1	4,085,897 2,314,336 8,255,684	1,777,929	2.910.290	135.5	6,016 518 2,126	296,985 80,300 430,893	1.9	165.564	570.229 162.572 829.715
Rhode Island South Carolina South Dakota	1,998,708 5,459,165 6,011,479	1,014,572	2,142,361 5,080,344 6,206,513	1,858,334 4,646,088 5,046,355	20%, 739 365,578 668,338	391.7	2,462,517 2,100,519	79.746	614.156 1.675.694 1.372.984	16.9 229.9 361.5	27.225	36.815 66.990 362,719	1.2 4.0 4.0	60.634 50.430 246.258	158.863 662.692 643.602
Tennessee Texas Utah	8,492,619 24,24,024 4,194,708	12,291,255	9,440,710 26,579,948 5,469,792	7.692.903	1,007,766 2,413,912 946,763	2,168.4	3.041.605 9.248.040 1.536.328	647.692 954.287 360.153	7.907.205	530.3	28,090 27,140	1,143,107	1.1	131,734 271,668 8,337	577.383 627.030 138.577
Vermont Virginia Washington	1,867,573	3,765,347	2.467.174 8.323.716 7.036.158	1.806.908 6.644.620 5.632.732	1,090,583	116.6 433.4 259.5	2,804.093	26,801	\$10.540 1.971.201 1.805.387	149.2	4,042 161.555	43.110 507.740 137.968	20.5	29,821	7,236 195,864 35,678
West Virginia Wisconsin Wyoming	4, 474, 234 9, 724, 661 4, 501, 327	2,280,335	4,249,980 10,718,068 5,125,126	3,738,906 9,052,904 4,123,788	387,201 1,184,352 817,355	146.5 460.6 719.2	1,655,705	671,325 532,063 363,973	3,179,755	133.1	56.511 82,664	142,561 408,632 84,735	2000	7,492 57,250 13,566	766,192 169,097 69.021
Destribt of Columbia	1,918,469	973,842	2,038,295	1,668,009	370,286	16.9	1,430,997	250,164	393.065	20.8	20,973	75.395	9.9	18,793	135.097
TOTALS	394,000,300	200,000,000	W216,155,312	345,064,988	W5.439.500	26,00%.8	170,845,754	41,523,397	114,215,941	7.860.9	2,516,501	17,531,538	876.3	4.895.014	22,613,021